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31 10:47:05 2005	GenCore version 5.1.6  Copyright (c) 1993 - 2005 Compugen Ltd.  c - nucleic search, using sw model  August 30, 2005, 04:38:43 ; Search time 18292 Seconds (without alignments)	11663.477 Million cell updates/sec 8core: 4403 1: Ltcggcacgaggcgcggttgaccgcggtggagctccagct 4403 table: IDENTITY_NUC Gapop 10.0 , Gapext 1.0 1: 4708233 segs, 24227607955 residues	<pre>1 number of hits satisfying chosen parameters: 9416466 mum DB seq length: 0 mum DB seq length: 2000000000 mum DB seq length: 2000000000 -processing: Minimum Match 00*</pre>	GenEmbl:*  1: 9b ba:* 2: 9b htg:* 3: 9b_in:* 4: 9b_oin:* 5: 9b_ov:* 6: 9b_pat:* 7: 9b_ph:* 9: 9b_pi:* 11: 9b_ets:* 11: 9b_ets:* 12: 9b_py:* 13: 9b_ui:* 14: 9b_ui:* 15: 9b_ui:* 16: 9b_ui:* 17: 9b_ui:* 18: 9b_ui:* 19: 9b_ui:	Score Match Length DB ID Description  4164.2 94.6 4406 9 AF058696 4102.4 93.5 4482 9 AF058696 4108.4 93.3 4486 9 AF0533599 4108.8 42.3 55134 9 AF0586291 4162.8 42.3 184919 9 AF068291 AF068291 Homo sapi BF0.8 42.3 31864 9 AF049895 Homo sapi AF068291 42.3 31864 9 AF049895 Homo sapi BF0.8 42.3 31864 9 AF049895 BF0.8 AF068291 Homo sapi BF0.8 42.3 31864 9 AF049895 BF0.8 AF069291 Homo sapi BF0.8 42.3 31864 9 AF049895 BF0.8 AF069291 Homo sapi BF0.8 42.3 31864 9 AF049895 BF0.8 AF069291 Homo sapi BF0.8 41.7 2044 6 AF085805 BF0.8 AF069291			
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KQI PKGKTP I FLARKQHKKLSSA VVPGGGERALI TEBRIBEERHPFLAFGTVVPTTT
NSGTLI PDCGKKWI OS IMDMLOROGLR PI PEAEI GLAVI FWITTKNY CDPQGHPSTGLK
TTTPGPSLSGGVS VDEKLMS BAPVYTTTYVADTES CAADTWDLSER PKE I KVSKMEQK
FRMLSQDAPTVKES CKTS SNNNSWYSNTLAKOR I PNYQLS PTKLES I KVSKMEQK
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ENES IS NNYPGPSE TKKRER NDEBNQENSS CKSAR I ETS CSLLEQTOPAT PSLWKNKEQHL
SENEPVDTNSDNNLFTDTDLKS I VKNSAS KSHAAEKLASNKKRENDDVA I EDE VLEQ
EKDTKPELE I DVKVQKQEBDVNVRKR PRMD I ETNDT F SDEAVPESS K I SQENE I GKKR
EKKRYKKYTYPGAGKE I SNDKLQDDS EMLPRKLLLITEFRS LVI KNSTSRNPSG I NDDYGQL
KNFKKKKYTYPGAGKL PHI I GGSDLI AHHARKNTELEBWLRQEMBVOQHAKEES LA
DDLERYNPYLKRRR
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7; Mismatches
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Best Local Similarity 97.1%;
Matches 4287; Conservative
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Mammalia; Butheria; Primates; Catarrhini; Hominidae; Homo.
Uases 1 to 4412)
Varon, R., Vissinga, C., Platzer, M., Cerosaletti, K.M.,
Chrzanowska, K.H., Saar, K., Beckmann, G., Seemanova, E., Cooper, P.R.,
Nowak, N.J., Stumm, Weemaes, C.M.R., Gatti, R.A., Wilson, R.K.,
Digweed, M., Rosenthal, A., Sperling, K., Concannon, P. and Reis, A.
Nibrin, a novel DNA double-strand break repair protein, is mutated
in Nilmegen breakage syndrome
Cell 93 (3), 467-476 (1998)
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Varon, R. and Platzer, M.
Direct Submission
Submitted (26-FEB-1998) Genome Analysis, Institute of Molecular Biotechhology, Beutenbergstr.11, Jena 07745, Germany
Location/Qualifiers
                                         AGCTATAATTGGGTCATAGAAATTCTTTATACATTCTAGATGCAAGTCTCTTGTTGTTA
                                                                                                                | TACGTAITGAGATAITACACCTAGTCTGTGGCTTGACTGTTTTCTTTAIGTCTTTTGATG
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/note="mutated in Nijmegen Breakage syndrome"
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Homo sapiens nibrin (NBS) mRNA, complete cds.
AF051334
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Db 4020 GTCATAGAAATTCTTATACATCTAGATGCTCTTGTCGGATATAGTATTGAGA 4079  Qy 4014 TATTACACCTAGTCTGTGGCTTGATCTTTATGTTTTGATGAATAGAATTTT 4039  Qy 4080 TATTACACCTAGTCTGTGGCTTGACTGTTTTTTTTTTTT	RESULT 3 CQ725600 CQ725600 CQ725600 DDA linear PAT 03-FEB-2004 DEFINITION Sequence 11534 from Patent W002068579. ACCESSION VERSION CQ725600. GI:42286781 KEYWORDS ORGANISM Homo sapiens (human) ORGANISM Homo sapiens (human) CG725600. GI:42286781 CG725600 CG72600 CG72600 CG72600 CG72600 CG72600 CG72600 CG72600 CG72600 CG726	TOURNAL Patent: WO 02068579-A 11534 06-SEP-2002;  PE Corporation (NY) (US)  FEATURES  Location/Qualifiers  1. 4388  Source  Location/Qualifiers  1. 4388;  Anol 1/Ppe="unassigned DNA"  Anol 1/Ppe="unassigned DNA"  Ab_xref="taxon:9606"  ORIGIN  Query Match  Best Local Smilarity 97.2%; Pred, No. 0; Tadala 91. Green of the control of the	COURSELVALIVE  COGNIGERATIVE  COGNIGERATIVE  COGNIGERATIVE  COGNIGERATOR COCCEGE  COGNIGERATOR COCCEGE  COGNIGERATOR COCCEGE  COGNIGERATOR COCCEGE  COGNIGERATOR COCTION  COCCEGE  COCC
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841 TACAGGAATAACAAACTCACAGACCTTAATTCCTGACTGA	1048 ATTAAAGACAACTCCAGGACCAAGCCTTTCACAAGGCGTGTCAGTTGATGAAAACT 1107			ATTCAGAATGCTTTCACAACCACCCACTGTAAGGAGCCTGCAAAACGAAACCTAAAAAGGAGCCTGCAAAAAGGAGCCCTGCAAAACCTAAACTGCAAAACGAAACCTAAAAAGGAGCCCTGCAAAAAAAA	TAATAATAGTATGGTATCAAJTACTTTGGCTAAGATCAGAATCCCAAACTATCAGCTTTC 	1348 ACCAACTAAATTGCCAAGTATAAATTAAAATTAAAGATAGGGCTTCTCAGCAGCAGCAGC 1407 	1408 CAACTCCATCAGAAACTACTTTCAGCCGTCTACCAAAAAAAGGGAAAGGGATGAAGAAA 1467 	1468 TCAAGAAATGTCTTCATGCAAATCAGCAAGAATAGAAACGTCTTGTTCTTTTAGAACA 1527 	1528 AACAACCTGCTACACCCTCATTGTGGAAAAATAAGGAGCAGCATCTATCT		GAAAATTCTGCCAGTAAATCTCATGCTGCAGAAAAGGTAAGATCAAJTAAAAAAGGGA 	AATGGATGATGTGGCCATAGAAGATGAAGTATTGGAACAGTTATTCAAGGACACAAAACC 	1768 AGAGTTAGAAATTGATGTGAAAAGTTCAAAACAGGAGGAAGATGTCAATGTTAGAAAAG 1827 	1828 GCCAAGGATGGATATAGAAACAAATGACACTTTCAGTGATGAAGCAGTACCAGAAAGTAG 1887 	1888 CAAAATATCTCAAGAAATGAAATTGGGAAGAAACGTGAACTCAAGGAAGACTCACTATG 1947 	1948 GTCAGCTAAAGAATATCTAACAATGACAAACTTCAGGATGATAGTGAGATGCTTCCAAA 2007 
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HORS Concannon, P.J., Vissinga, C.S., Cerosaletti, K.M., Varon-Mateeva, R., Sperling, K. and Reis, A.W. S.  LE Gene associated with Nijmegen breakage syndrome, it's gene product and methods for their use and methods for their use and methods for their use Location/Qualifiers  RES Location/Qualifiers  1. 4386  / organism="unknown"  / mol_type="genomic DNA"	584	28 GCCCAGCCCTGAGGAGCGGACCGATGTGGAAACTGCTGCCGCGGGGGGCCCGGCAGG 87	88 AGGAGAACCATACAGACTTTTGACTGGGTTGAGTACGTTGTGGAAGGAA	148 CATTCTAATTGAAAATGATCGATCAGCCGAAATCATGCTGTGTTAACTGCTAACTT 207 	208 TICTGTAACCAACCTGAGTCAACAGATGAAATCCCTGTATTGACATTAAAAGATAATTC 267 	268 TAAGTATGGTACCTTTGTTAATGAGAAAAAATGCAGAATGGCTTTTCCCGAACTTTGAA 327 	328 GTCGGGGGTATTACTTTTGGAGTGTTTGGAAGTAAATTCAGAATAGAGTATGACC 387 	TITGGTTGCATGCTCTTCTTGTTTAGATGTCTCTGGGAAAACTGCTTTAAATCAAGCTAT	GTCAT	ATTGT        ATTGT	568 AAAGCCAGAATATTTACTGAATTCCTGAAAGCAGTTCAGTCCAAGAAGCAGCTCCACA 627 	628 AATTGAAAGTTTTTACCCACCTCTTGATGAACCATCTATTGGAAGTAAAAATGTTGATCT 687 	688 GTCAGGACGGCAGGAAAGAAATCTTCAAAGGAAAACATTATATTTTTGAATGC 747 		808 AACAGAAGAGAATGAAGAACATAATTTCTTTTTGGCTCCGGGAACGTGTGTTGA 867 	TACAGGAATAACAAACTCACAGACCTTAATTCCTGACTGTCAGAAGAAATGGATTCAGTC
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                                                     Clone distribution: MGC clone distribution information can be foun through the I.M.A.G.E. Consortium/LLNL at: http://image.llnl.gov Series: IRAK Plate: 168 Row: m Column: 13
This clone was selected for full length sequencing because it passed the following selection criteria: matched mRNA gi: 6996019.
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   Dickson, M., Schmutz, J., Grimwood, J., Rodriguez, A., and Myers,
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7; Mismatches
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Homo sapiens Nijmegen breakage syndrome 1 (nibrin), mRNA (cDNA
clone MGC:87362 IMAGE:30343504), complete cds.
BC071590.1 GI:47938122
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DNA Sequencing by: Sequencing Group at the Stanford Human Genome
Center, Stanford University School of Medicine, Stanford, CA 94305
Web site:
                 CAAATTTATTTTTTTTTTTTTTTTTTTTTTTTTCCCCAATTTTAACCCCAAGATTTCA 4188
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Submitted (01-JUN-2004) National Institutes of Health, Mammalian
Gene Collection (MGC), Cancer Genomics Office, National Cancer
Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590,
                                                                                                                                                                                                      GATATTCTGCTCTATTATATAAACTTTATATTTTTTATATTTGTGATCTACCTTGAATTGA
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                                                                                                                                                               4249 TATGTATGTTGTGAATTATGGATCAGGGTTCTTTTTTTCCCCCATACAAGTATCCAGTCA
Email: cgapbs-remail.nih.gov
Tissue Procurement: Dr. Stefan Hansson
cDNA Library Preparation: Michael Brownstein / Ted Usdin
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Contact: MGC help desk
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PUBMED
REFERENCE
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                                                                                                                                                                                                             Gaps
                                                                                                                                                                                     91.7%; Score 4038.4; DB 11; Length 4423;
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Conservative 8; Mismatches 41; Indels 103;
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submitted
    ð
   Diego,
  3595 John Hopkins Court, San Di
Tel: 18882029018
Fax: 18582029020
Email: abraun@sequenom.com
Primer A: No primer sequence su
Primer B: No primer sequence su
STS size: 4423.
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Matches 4270; Conserv
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repeat_region variation variation	variation	variation variation variation	variation gene mRNA	CDS			variation	variation
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	56500 k or NBS1, cc 124 cage syndrc	Eukan Mamme 1 1 Matsu Endo, Oshir Posit Nat.	PUBMED 9620777  REFERENCE 2 (bases 1 to 56500)  AUTHORS Matsuura, S., Tauchi, H. and Komatsu, K.  AUTHORS Matsuura, S., Tauchi, H. and Komatsu, K.  TITLE Direct Submission JOURNAL Submitted (21-APR-1998) Shinya Matsuura, Hiroshima University, Research Institute for Radiation Biology and Madicine, Department of Radiation Biology, Kasumi 1-2-3, Minami-ku, Hiroshima, Hiroshima 734-8553, Japan (B-mail:shinya@ue.ipc.hiroshima-u.ac.jp,	Tel:81-82-257-5811, Fax:81-82-256-7101)  COMMENT Sequence updated (26-May-1998).  Location/Qualifiers  156500  /mol_type="genomic DNA"	/db xref="taxon:9606" /chromosome="8" /map="8q21" /map="8q21" /sub_clone="RG255A7" n 4790. 4848 /number=1 /rt 4790. 4811	CDS join (4812. 4848, 6518. 6651, 7850. 7998, 8480. 8639, 11050. 11153, 18083. 18200, 18816. 19009, 24866. 24963, 30519. 30648, 33818. 34090, 35682. 36129, 41481. 41549, 43078. 43233, 46007. 46120, 52298. 52347, 53761. 53791) /codon_start=1 /product="NBS1" /product="RBS1" /product="RBA28616.1" /product="RBA28616.1"	/ translation="MWKLLPAAGPAGGPPYRLLTGVEYVVGRKNCAILIENDQSISRN HAVLTANFSVTNLSQTDEIPVLTLKDNSKYGTFVNBEKMQNGFSRTLKSGOGITFGVF GSKRRIETSPELVACSSCLDVSGKTALNQAILOLGGFTVNNWTEBCTHLWWSVKYTIK TICALIGRPIVKPSYFTBETLAVBSKKQPPQISSFYPPLDEPSIGSKWVDLSGRQBR KQIFKGKTPFFLUNKQHKKLSSAVVFGGGBRALLTERNBEBHNFFLAPGTCVVDTGIT NSQTLIPDCQKKWIQSIMDMLQRQGIRPIPBABIGLAVIFWTTKNYCDPQGHPSTGLK TTTPGPSLSQGVSTDEKLMPSPANYTTYNADTSESBADTWALTKNYCDPQGHPSTGLK GDALFONYGGGGRAPHTTANYCDPSTGRANDAILSENSKEIKSKRENSKENSKEIKSK	OTNGI RNY POSTKRERDEENQEMSSCKSAR I ETSCSLLEQTQPATPSLWKNKEQHL SERREYDTNSDNILFTDTDLKSI VKNSASKSHAAELKESKKKERDDVALEDDFLEQL FKOTKPELEI DVKVQKQEEDVNVRKR PRID I ETNDTFSDEAVPESSKI SQENEI CEKKR ELKEDSLWSAKEISNNULSKLQDDSRMLLFKGLLLTERSLVI KNSTSRNPSGINDDYGQL KNFKKFKKVTYPGAGKLPH I I GGSDLI AHHARKNTELEEMLRQEMEVQNQHAKEESLLA DDLFRYNPYLKRRR"
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/number=3 7363573826 /gene="hT41"	/ number=4 80160 80251 /gene="hT41" /number=6	0.358386947		for Nijmegen breakage sy. join(9451294692,96106. 5375,106904106972,1123	1.14555. 11/1002311/3946. 1124590122587, 129444129637,130253130370,137300137403. 139814139973,140455140603,141802141935,	14360514363))   Jeene="NBS1"   / Jeene="NBS1"	complement (9451294692)	/unuberard /unuberard /ounplement(join(94662. 94692,96106. 96155,102333102446, 105220105375,106904106972,112324112771,	114363114635,117805117934,122490123587, 129444129637,130253130370,137300137403, 139814139973,140455140603,141802141935, 143605143641)	/gene="NBS1" /note="similar to yeast Xrs2" /codon_start=1	/product="nibrin" /protein_id="AAC6223.1"	/db_xref="G1:3687830" /tdb_xref="G1:3687830" /tdb_xref="G1:3687830"	GSKFILEVEVILLEGVILLEGNSKULF VREEKRIGNESKLUIFGVE GSKFILEVEPLVACSSCLDVSGKYALNQALLQLGGFTVNNWTEECTHLVWVSVKVTIK TICALIGGRPIVKPETFELKAVESKKQPPQIESRYPPIDEPSIGSKNVDLSGRQFR VOITKVGWTFTENNAKOUVUT SCANTEGGERBITTREPRINDERINDERINDERINDERINDERINDERINDERINDE	NSQTLIPCQKKWIQSIMMLQRQSLRPIPEAEIGLAVIFWITKWYCDPQGHPSTGLK TTTPGPSLSQGVSVDEKLMPSAPVNTTTYVADTESEQADTWDLSERPKEIKVSKMEQK	FRMLSQDAPTVKESCKTSSNNNSNVSNTLAKMRI PNYQLSPTKLPSINKSKDRASQQQ QTNSIRNYFOPSTKKRERDEENOEMSSCKSARIETSCSLLEOTOPATPSLMKNKEOHL	SENEPVDTNSDNNLFTDTDLKSIVKNSASKSHAAEKLRSNKKREMDDVAIEDEVLEQL FKDTKPELEIDVKVQKQEEDVNVRKRPRMDIETNDTFSDEAVPESSKISQENEIGKKR	ELKEDSLWSAKEISNNDKLQDDSEMLPKKLLLTEFRSLVIKNSTSRNPSGINDDYGQL KNFKKFKVTYPGAGKLPHIIGGSDLIAHHARKNTELEEWLRQEMEVQNQHAKEESLA DDLFRYNPYLKRRR"	complement(9610696155) /gene="NBS1" /grammer="1"	/luminet 1.0 complement (102333102446)	/ Journal   / Jour	/number=13 _complement(106904106972)	/gene="NBS1" /umber=12 /complement(112224 112221)		complement (114363114635) /gene="NBS1"	/number=10 complement(117805117934) / gene="NBS1"	/number=9 complement(123490123587) /gene="NBS1"
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                                                                                                                                                                                                                                                                                                                                                                  Neighboring sequence information:
This entry is part of a larger genomic contig. The start of this sequence is directed towards the centromere. The start of (1. .2000) overlaps with the end of the neighbouring Acc number AF117829. The end (329871. .331870) of this sequence overlaps with the start of Acc number AF17830. It is overlapped by SCb-296N11, SCb-216M22, SCb-296C9 and covers SCb-284N21, SCb-157K21, SCb-58110, SCb-228C20 entirely.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     regions were double stranded, sequenced with an alternate chemistry, or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by at least one plasmid subclone or more than one M13 subclone; and the assembly was confirmed by restriction digest.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                This sequence was finished as follows unless otherwise noted: all
                                                                                                                                                                                                                                           Direct Submission
Submitted (04-NOV-2003) Genome Analysis, Institute of Molecular
Submitted (04-NOV-2003) Genome Analysis, Institute of Molecular
Biotechnology, Beutenbergstr. 11, Jena, Thuringia 07745, Germany
4 (bases 1 to 311844)
Lagemann, D. and Platzer, M.
Direct Submission
                                                                  2 (bases 1 to 331864)
Platezer,M. and Varon,R.
Direct Submission
Submitted (30-DEC-1998) Genome Analysis, Institute of Molecular Blotecenhology, Beutenbergstrasse 11, Jena 07745, Germany
Sequence update by submitter
(bases 1 to 331864)
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        Sequence Quality Assessment:
This entry has been annotated with sequence quality estimates computed by the Phrap assembly program.
All manually edited bases have been reduced to quality zero. Quality levels above 40 are expected to have less than 1 error in 10,000 bp.
Base-by-base quality values are not generally visible from the GenBank flat file format but are available as part of this entry's ASN:1 file.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            This clones were finished using overlapping sequence from accessions AC004083, AC123779, AC004612
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            Assembly program: Phrap; version 0.990329
Consensus quality: 326110 bases at least Q40
Consensus quality: 330377 bases at least Q30
Consensus quality: 331864 bases at least Q20
Quality coverage: 6.80x
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SOURCE INFORMATION:

This clone was derived from human PAC library RFCI-5, prepared by Pieter de Jong and coworkers at the Roswell Park Cancer Institute (http://bacpac.med.buffalo.edu) using the method described by Ioannou et al., Nature Genetics 6:84-9 (1994). The library is from one male donor.

The clone may be obtained either from Genome Systems, Inc.

(http://www.resgen.com); or from Pieter de Jong.

VECTOR: pCYPAC.

NEIGHBORING SEQUENCE INFORMATION:

The clone sequenced to the left is CTA-437L15, 200 bp overlap; the clone sequenced to the right is RPI-31847, 200 bp overlap; Actual start of this clone is at base position 115721 of CTA-437L15; actual end is at base position 11522 of RPI-31847.
               Louis
                                                                                                                                                                                                                                                                                       NOTICE: This sequence may not represent the entire insert of this close. It may be shorter because we only sequence overlapping clone sections once, or longer because we provide a small overlap between neighboring data submissions.
                                                                                                                                                                                                                                                                                                                                                                                          This sequence was finished as follows unless otherwise noted: all regions were double stranded, sequenced with an alternate chemistry, or covered by high quality data (i.e., phred quality 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by sequence from more than one subclone; and the assembly was confirmed by
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Mapping information for this clone was provided by Dr. John D. McPherson, Department of Genetics, Washington University, St. I. MO. For additional information about the map position of this sequence, see http://genome.wustl.edu/gsc
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4 (Dases 1 to 107549)
Waterston, Waterston, Dabirect Submission
Direct Submission
Submitted (14-OCT-2000) Department of Genetics, Washington
University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
                                                                                                                                                                                                                                                   3998
                                                                                                                                                                                                                                                                                       4990
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Homo sapiens PAC clone RP5-1098020 from 8, complete sequence.
ACO74178
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4930
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Sulston, J. B. and Waterston, R. Toward a complete human genome sequence
                                                                                                                                                                                                                                                                        GATAGCTATAATTGGGTCATAGAAATTCTTTATACATTCTAGATGCAAGTCTCTTGTCGG
                                                                                                                                                                                                                                                                                                                                                                                                                                     5104 CCATTCCAATCGGTGTGTGTGTG----TITCATTTTGGTTTTAATTTGTATATCCCT
                                                                                                                                                                                                                                              GATAGCTATAATTGGGTCATAGAAATTCTTATACATTCTAGATGCAAGTCTTTGYCGG
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Submitted (15-JUL-2000) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St.
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Unpublished
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Homo sapiens (human)
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Dante, M.
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/rpt_family="L1" repeat_region 2179321917	repeat_region 219182210 / rrr family="Alu"	repeat_region /22/12380 /22112380 /rot family="A]u"			repeat_region 2361723881 /rpt_family="Alu"		24697 /rpt	~		repeat_region 2967731685 /rpt_family="L1"	egion 3168 <u>9</u> 31818	Query Match 42.3%; Score 1861.2; DB 9; Length 107549; Best Local Similarity 94.1%; Pred: No. 0; Matches 2023: Conservative 9. Mismatches 27: Indels 91: Gans 6:	Matches 2023; Competivative 3; Manuaca 27; Matches 27; Cabe	ZZ/8 ICITITITAGATACATICCTTATITAAAAGAAGAAGAAGAACITAAGAAGAAGAAGAAGAAGAAGAAGAAGAAGAAGAAGAAG	68939 TCTTTGCAGATACCATTTTTAAAAAGGAAGATAACTGAGGATTTTAAAAAGAAG	Oy 2338 CCATGGAAAACTTCCTAGTAAGCATCTACTTCAGGCCAACAAGGTTATATGAATATATA 2397	000/) CONTOCERANTANCE INCLINATIONAL CINCLICANCE AND CONTOCERATE AND CONTOCERAT	6,0819 GTGTTATAGAGTGTATAGAGTTTAGAGTGTTAGAGTGTTAGAGTGTTAGAGTGTAGAGTGTAGAGTGTGTTAGAGTGTGTGTAGAGT	2458 CAAAACTITTICATTITICATIONAACTAACTAATTICATTICATYCTYCTTCAGGCTTTTICATCATTIC	68759 CAAAACTTTGATTCTTTTGTAACAATTGTTGTTCTGTTTTCAGGCTTTTGTCATTG	Qy 2518 CATCTTTTTCATTTTTAAATGTGTTTTTTAAATAGTTAATAATATAGTCACAGTTC 2577	68699 CATCTTTTTTCATTTTAAATGTGTTTTTGTTTAAATAGTTAATAATAATTAAGTCACAGTTC	Qy 2578 AAAATTCTAAATKTACGTAAAGGTAAAGGACTAAAGGTCACCCTTCCACCATTCTCCTAGCT 2637	Db 68639 AAAATTCTAAATACGTAAGGTAAA-GACTAAAGTCACCCTTCCACCATTGTCCTAGCT 68581	Qy 2638 ACT	   68580 ACTTGGTTCCCCTCAGAAAAATTCATGATACTCATTTCTTATGAATCTTTCCAGGGAT 68521	Qy 26412678	Db 68520 TTTTGAGTCCTATTCAAATTCCTATTTTTAAATAATTTCCTACAAATGATAGATA	Oy 2679 ATATGCAGTGTTCTACATTGCTTTTTTACTTAGTAAGATTAAAAATTATAGGAATATC 2738	68460 AIAIGCAGIGITCTACACCTTGCTTTTTTACTTAGT-AGATTAAAAATTATAGAAATAC	Ov 2739 AATATAATGTTTTTAATATTTTTTCTTTTCCATTATGCTGTAGTCTTACCTAAACTCTGG 2798	68401 AATATAATGTTTTTAATATTTTTTCTTTTCCATTATGCTGTACTCTTACCTAAACTCTGG	2799	Db 68341 TGATCCAAACAAAATGGCTTCAGTGGTGCAGATGTCACCTACATGTTATTCTAGTACTAG 68282	
e 21272549 /note="similar to EST N66848 (NID:g1218973) za46d10.s1"		/rpt 4114		/rpt_family="Alu" on 4289, 4425	/rpt_family="Alu" 44304509	/rpt_family="MER1 45465169			/rpt_family="Alu" 60146289																									on 20787, .20974 /rpt_family="L1"	
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2859 AAACTGAAGACCATGTGGAGACTTCATCAACATGGGTTTTAGTTTTCACCAGAATGGAAA 2918 68281 AAACTGAAGACCATGTGGAGACTTCATCAACATGGGTTTAGTTTTCACCAGAATGGAAA 68222 2919 GACCTGTACCCCTTTTTGGTGGTCTTACTGAGCTGGGTGGG	3279 GCCTCTACATTACTTCCAATCTTGGGAAGTGCATCTACTTGCCAACCAA
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1153   AGANTCAGGOARGCAATCACTGCAATTCAGGGCAAACCAAAAAACCTGCTTC   1262	RESULT 13 AX875862 LOCUS LOCUS DEFINITION Sequence 10767 from Patent EP1074617. ACCESSION AX875862 VERSION XX875862.1 GI:40030598
6) COCCECTION COCCECTORACCION COCCECTORACTICACION COCCECTORACTICACION COCCECCOCOCCECACION COCCECCOCOCCECCOCCECCOCCECCOCCECCOCCECCOCCECCOCCECCOCCECCOCCECCOCCECCOCCECCCOCCECCCOCCECCCCCC	1033 CCATCCCAGTACAGGATTAAAGACAACAACTCCAGGACCAAGCCTTTCACAAGGCGTGTC 1092

	1027 AGCAGAAATTGGATTGGCGGTGATTTTCATGACTACAAAGAATTACTGTGATCTCTGAGGG 1086 1027 AGCAGAAATTGGATTGGCGGTGATTTTCATGACTACAAAGAATTACTGTGATCCTCAGGG 1086 1033 CCATCCCAGTACAGGATTAAAGACAACAACTCCAGGACCAAGCCTTTCACAAGGCGTGTC 1092 1087 CCATCCCAGTACAGGATTAAAGACAACACCAGGACCAAGGCCTTTCACAAGGCGTGTC 1146	1093   AGTTGATGAAAACTAATGCCAAGCGCCCCAGTGAACATACAT	1213 CAAAATGGAACAAAAATTCAGAATGCTTTCACAAGACGCACCCACTGTAAAGGAGTCCTG	1333   AAACTATCAGCTTTCACCAACTAAATTGCCAAGTATAAATAA	1452 AAAGGGATGAAGAAATGTCTTCATGCAAATCAGCAAGAATAGAAACGTCTT   1511
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SOURCE Homo sapiens (human) ORGANISM Homo sapiens UREFERENCE Homo sapiens  AUTHORS Ishii,S. Sugiyama,T., Nishikawa,T., Hayashi,K., Saito,K., Yamamoto,J., TITLE Primers for synthesising full-length cDNA and their use JOURNAL Research Association for Biotechnology (JP)  FRATURES  Location/Qualifiers Source /organism="Homo sapiens" /db xref="taxon:9606" /db xref="taxon:9606" /db xref="taxon:9606" /codon start=1 /protein_id="CAE89934.1" /db xref="di:40030599" /translation="WMKLLPAAGPAGREPYRLLTGVEYVVGRKNCAILIENDGSISRN HAVLTANFSVTNLSQTDEIPVLTLKDNSKYGFFVNDESCHUNNSKYRTIK TICALICGREIVKEBSFFTEREKANOSKYGPFVNDESCHUNDSGREN  KOIFKGKFFFILMSQTMETELASSAVVFGGGEARLITTEENSEEHNFILAPGTCVVDTGIT	ORIGIN  Query Match  41.7%; Score 1836.4; DB 6; Length 2044;  Best Local Similarity 94.3%; Pred. No. 0;		127 CGCGGGCCCGGCAGCAGCACATACAGACTTTTGACTGGCGTTGAGTACGTTGTTGG 18  133 AAGGAAAACTGTGCCATTCTAATTGAAAATGATCAGTCGATCAGCCGAAATCATGCTGT 19  187 AAGGAAAAACTGTGCCATTCTAATTGAAAATGATCAGTCGATCAGCCGAAATCATGTGT 24  193 GTTAACTGCTAACTTTCTGTAACCAACCTGAGTCAAACAGATGAATCCCTGTATTGAC 25	247 GTTAACTGCTAACTTTTCTGTAACCAGCTGAGTCAAACAGATGAATCCCTGTATTGAC 30 253 ATTAAAAGATAATTCTAAGTALGGTACCTTTGTTAATGAGGAAAAAATGCAGAATGGCTT 31 307 ATTAAAAGATAATTCTAAGTATGGTACCTTTGTTAATGAGGAAAAATGCAGAATGGCTT 36 313 TTCCCGAACTTTGAAGTCGCGCACGATGATTACTTTTCGAGTGTTTGGAAGTAAATTCAG 37	Db   367   TTCCCGAACTTTGAAGTCGGGGGGATGGTATTTTGGGGGTTTTGGAAGTAAATTCAG   426

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Jacogal, T. and Otsuki, T.

Submitted (16-FEB-2000) Takao Isogai, Helix Research Institute,
Submitted (16-FEB-2000) Takao Isogai, Helix Research Institute,
Genomics Laboratory; 1532-3 Yana, Kisarazu, Chiba 292-0812, Japan
(E-mail:genomics@hir.co.jp, Tel:81-438-52-3375, Fax:81-438-52-3986)
NEDO human CDNA sequencing project supported by Ministry of
International Trade and Industry of Japan, CDNA full insert
sequencing: Research Association for Biotechnology; CDNA library
construction, 5' & 3'-end one pass sequencing and clone selection:
Helix Research Institute (supported by Japan Key Technology Center
etc.) and Department of Virology, Institute of Medical Science,
                                           Isogai, T., Ota, T., Hayashi, K., Sugiyama, T., Otsuki, T., Suzuki, Y., Nishikawa, T., Nagai, K., Sato, H., Sugano, S., Shiratori, A., Sudo, H., Wagatsuma, M., Hosoiri, T., Kaku, Y., Kodaira, H., Kondo, H., Ono, Y., Takiguchi, M., Chiba, Y., Ishida, S., Murakawa, K., Ishii, S., Yawai, Y., Saito, K., Yamamoto, J., Wakamatsu, K., Nishii, S., Kawai, Y., Saito, K., Yamamoto, J., Wakamatsu, A., Nakamura, Y., Nagahari, K., Masuho, Y., Ninomiya, K. and Iwayanagi, T. NEDO human cDNA sequencing project
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/replace=""
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    /tissue_type="whole embryo, mainly head"
/clone_lib="HbmBA1"
/dev stage="embryo.10 weeks"
/note="cloning vector: pME185FL3"
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Pred. No. 0;
0; Mismatches
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/mol_type="mRNA"
/db_xref="taxon:9606"
                  36 (1), 40-45 (2004)
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94.3%;
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Matches 1976; Conservative
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2044 bp mRNA linear FRI 30-JAN-2004
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NAMA Sapiens gene for NBS1.
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                                                                                                       /note="contains forkhead-associated (FHA) and bream cancer carboxy terminal (BRCT) domains; mutated in Nijmegen breakage syndrome" /codon_start=1
                                                            RAD50 and NBS1
in response to DNA
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Pred. No. 3e-243;
0; Mismatches 546; Indels
                                                          /function="required for MRE11,
colocalization at nuclear foci
double-strand breaks"
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76.1%;
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Search completed: August 30, 2005, 11:06:46 Job time : 18315 secs

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Human gen Bovine ES Human imm

Adp72176 Renal tox

OM nucleic

Run on:

Sequence:

Searched:

Database

Result No.

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The invention relates to a method for the detection of cancer in which a change in the expression of 1 or more of 264 specified cancer associated genes, ABZ71694-ABZ71957, or of sequences at least 80% homologous to them in the specimen tissue as compared to normal tissue is observed. The genes are used in detection, diagnosis and treatment of cancer, associated polymucleotide of the invention. Note: The present sequence was not given in the printed specification but was isolated using the
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               Measuring changes in expression of 264 cancer associated genes for detection of stomach cancer and screening of potential anticancer agents.
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                     CATTCTGATTGAAAATGATCAGTCGATCAGCCGAAATCATGCTGTTAACTGCTAACTT
                                                                            TTCTGTAACCAACCTGAGTCAAACAGATGAAATCCCTGTATTGACATTAAAAGATACTTC
                                                                                                                                      TAAGTATGGTACCTTTGTTAATGAGGAAAAATGCAGAATGGCTTTTCCCGAACTTTGAA
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                                                                                       ACAAGTATCCAGTCATTGTAACACTGTTTATTGAAAGAATTATCCTTTCCTCATTAAATT
 TCTACCTTGAATTGATATGTTGTGAATTATGGATCAGGGTTCTTTTTTTCCCCCAT
                                                           ACAAGTATCCAGTCATTGTAACACTGTTTATTGAAAGAATTATCCTTTCCTCATTAAATT
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                                                                                                                                                                                                                                                                                                                                               Nibrin; human; DNA double strand break repair protein; diagnosis; therapy; Nijmegen Breakage Syndrome; gene therapy; ds.
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<i>장</i> 옵	1288 TAATAATAGTATGGTATCAATACTTTGGCTAAGATGAGAATCCCAAACTATCAGCTTTC 1347 	<u></u>	TTCAGGCCAACAAGGTTATATGAATATATATATGTATAGAAGCGATTTAGGTTACAATGTT
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ે જે જે	CAACTCCATCAGAAACTACTTTCAGCGTCTACCAAAAAAAGGGAAAGGGATGAAGAAAAAAAA	& A	2488 TGTTTGTYCTGTTTTCAGGCTTTGTCATTGCATCTTTTTTCATTTTTAAATGTGTTTTG 2547 
3 & 8	TCAGAAATGTCTTCATGCAATCAGCAAGAATGAAACGTCTTGTTCTCTTTTAGAACATCAGAAAAGATCTTGTTCTTTTTTAGAACA	ò a	2548 TTTATTAAATAGTTAATATAGTCACAGTTCAAAATTCTAAATRTACGTAAGGTAAAGGAC 2607 
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<i>∂</i> 8	TATTGT 	<i>ò</i> 8	2641TATTTTA 2648 2640 TACTCATTCTTATGAATCTTTCCAGGGATTTTTGAGTCCTATTCAAATTCCTATTTTA 2699
8 6	GAAAATTCTGCCAGTAAATCTCATGCTGCAGAAAAGGTAAGGTCAATTAAAAAAGGGA 	ò a	2649 AATAATTICCTACACAAATGATAGCATAACATAIGCAGIGTICTACACCTIGCTITITA 2708 
è	AATGGATGATGGCCCATAGAAGATGAAGTATTGGAACAGTTATTCAAGGACACAAAACC	<i>&amp;</i> ₽	2709 CTTAGTAAGATTAAAGGAATATCAATATAATGTTTTTAATATTTTTTTCTTTC
}	AGAGTTAGAAATTGATGTGAAAAGTTCAAAAACGGGAGGAAGATGTCAATGTTAGAAAAG 182 AGAGTTAGAAATTGATGTGAAAGTTCAAAAACGGGAGGAAGAAGTTCAATGTTAGAAAAG 182 AGAGTTAGAAATTGATGTGAAAATTGATTCAAAAAAAAAGAAGAAAAATGTTAGAAAAAG 180	<i>&amp;</i> 8	2769 CATTATGCTGTAGTCTTACCTAAACTCTGGTGATCCAAAAAAAGGCTTCAGTGGTGCA 2828 
ે કે કે	CONSTRUCTION OF THE STATE OF TH	å a	2829 GATGTCACCTACATGTTATTCTAGTACTAGAAACTGAAGACCATGTGGAGACTTCATCAA 2888 
ි රි සි	CCAAAATATCTCAAGAAATGAAATTGGGAAGAAACGTGAACTCAAGGAAGAAGATTGGGAAGAAACGTGAACTCAAGGAAGAAGAACTTGGGAAGAAACTGAAATTGGGAAGAAACTGAAATTGGGAAGAAACTGAACTGAACTTGAAATTGAAATTGGGAAGAAACTGAACTGAACTTAATGAAATTAAGAAAATGAAAATGAAATGAAATGAAATGAAATGAAATGAAATGAAAATGAAATGAAATGAAATGAAATGAAAATGAAAATGAAAATGAAAATGAAAATGAAATGAAAATGAAAATGAAAATGAAAATGAAAATGAAAATGAAAAATGAAAATGAAAATGAAAATGAAAAAA	ò 8	2889 ACATGGGTTTAGTTTTCACCAGAATGGAAAGACCTGTACCCCTTTTTGGTGGTCTTAGTG 2948 
6 6	GTCAGCTAAAGAAATATCTAACAATGACAAACTTCAGGATGATAGTGAGATGCTTCCAAA  [	<i>≿</i> 8	2949 AGCTGGGTGTGTCTGTTTTGAGCTTATTTAGAGTCCTAGTTTTCCTACTTATAAAGTA 3008 
3 8 8	SCHOOLINGARING TO THE CONTROL OF THE	& 8	3009 GAAATGGTGAGATTGTTTTCTTTTCTACCKTAAGGGGATGGTAAGAAACAATGAATG 3068 
i & f	GTCTGGCATAAATGATTATGGTCAACTAAAAATTTCAAGAATTCAAAAAAGGTCACTACTACTAAAAAGGTCACTACTAAAAAAGATTCAAAAAAGGTCACTACTAAAAAAGTTTCAAGAAAAGGTCACTAAAAAAATTTCAAGAAAAAAGTTCAAAAAAAGGTCACTAAAAAAATTTCAAGAAAAAAAA	ò 8	3069 TCTTTTTCAAACTTTATTGACAAGTGATTTTCAAGTCTGTGTTCAAAAATATATTCATG 3128 
	ATAICCTGGAGCAGGAAAACTTCCACATCATTGGAGGATCAGATCTAAAAAGTCACATCATCATTGGAGGATCAGATCTAATAGCTCATCATTGGAGGATCAGATCTAATAGCTCATCATTAGAAGATCATCATTGAATAGTTCATAATAGTTCATCATCATCATCATCATCATCATCATCATCATCATCA	ò 8	3129 TACCTGTGATCCAGCAAGAAGGGGTTCCAGTCAAGAGTCACTACAACTGATTAGTTGTT 3188 
3 8 8	TGCTCGAAAGAATACAGAACTICCACAATGATGAAGGGAGAAATGGAAGATGAAATGAA	& 8 ———	3189 TAGAGAATGAGAAATGGAACAGTGAGGAATGGAGGCCATATTTCCATGACTTCCCTTGTA 3248 
ે જે સ	1GCICGAAAGAAIACAGAACIAGAAGAAGAGIGGCIAAGGGAAGAIGGAAGAIACAAGAACA 222 ACATGCAAAAGAAGAAGATCTCTTGCTGATGATCTTTTTAGATACAATCCTTATTTAAAAAG 230 ACATGCAAAAGAAGAAGATGTCTTTTTTTTTTTTTTTTT	& 8	3249 AACAGAAGCAACAGAAGAGACAAGAGCTGGCCTCTACATCACTCTCACCTTCCAAATCT 3308 
g &	2221 ACATGCAAAGAAGAGTCTCTTGCTGATGATCTTTTTAGATACAATCCTTATTTAAAAAG 2280 2308 GAGAAGATAACTGAGGATTTTAAAAAGAAGCCATGGAAAAACTTCCTAGTAAGCATCTAC 2367 	장 업	3309 TGTGGAAGTGCATCTACTTGCCAGAACTAAATTAACTTACTT

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                        3419 CAGGTGGAACTCCAGCTGCAAGGGAGTTAGGGGAAATGAAGGTCTTTTTTTAAAAGCTTCT
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This is the nucleotide sequence of cDNA for the NBS1 gene that is associated with the Nijmegen breakage syndrome (NBS). It includes a coding region for a 754-amino acid protein, inbrin (see AAY373). The coding region for a 754-amino acid protein, inbrin (see AAY373). The gene maps to a 1 cM region on chromosome 8021. The invention is based on the discovery that the gene contains mutations in all NBS patients. These mutations include deletions and insertions in all NBS patients. These well as point mutations. Specific mutations associated with the NBS phenotype include 657dels, 69ddel4, 842inst, 1142delC, 976CST, 681delT and 900del25. Polymorphisms include 553 G/C, 1197 T/C, 2016 A/G, 102 G/A, IVS 5+9 T/C, IVS5+51delT, IVS9+18 C/T and IVS-7A/G. It is an cobject of the invention to detect a mutation or pathological condition such as cancer, microcephaly, mental retardation, and primary ovarian failure, based on detection of a mutation in the NBS1 gene. It is also an object of the invention to treat NBS by replacing the mutated gene in a NBS patient by gene therapy. Recombinant vectors, genetically engineered host calls, a method for producing nibrin polypeptide, an antibody that specifically binds to the polypeptide, and method for antibody that specifically binds to the polypeptide, and method for antibody that specifically binds to the polypeptide, and method for antibody that specifically binds to the polypeptide, and method for antibody that specifically binds to the polypeptide, and method for producing number of the polypeptide.
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                                                    NBS1 gene; nibrin; Nijmegen breakage syndrome; diagnosis; human; gene therapy; cancer; microcephaly; mental retardation; primary ovarian failure; ss.
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gene associated with Nijmegen breakage syndrome
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7; Mismatches
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Best Local Similarity 97.1%;
Matches 4269; Conservative
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P-PSDB; AAY32373.
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AAZ34997 standard; cDNA; 4386 BP.

RESULT 3
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 This invention describes a novel DNA double strand break repair protein, Nibrin. Nibrin and DNA encoding it are useful for diagnosis and/or therapy of diseases influenced by repair of DNA-double strand breaks, in particular Nijmegen Breakage Syndrome. The product of the invention has applications in gene therapy. This sequence encodes the nibrin protein described in the invention
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                                                                                                                                                                                                                                                                                                                                                                                                    A DNA double strand break repair protein, Nibrin, and related DNA for diagnosis and therapy of Nijmegen Breakage Syndrome and other diseases influenced by DNA-double-strand break repair.
                                                                                                                                                                           diagnosis;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     91;
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                                                                                                                                                                      Nibrin, human; DNA double strand break repair protein; therapy; Nijmegen Breakage Syndrome; gene therapy; ds.
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9; Mismatches
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Best Local Similarity 94.1%;
Matches 2024; Conservative
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                                                                 DNA;
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                                                               standard;
                                                                                                                                             Human nibrin DNA
                                                                                                                                                                                                                                        DE19818680-C1
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                                                                                                                                                                                               ATATATGTATTGAGATATTACACCTAGTCTGTGGCTTGACTGTTTCTTTATGTCTTTTG
GCCATGTATCATAATTACCAAGTGAAGCTGGAACATATGGTCTCCCATTTTACAGTTA
                                                                                TITICICCCCAATITAACCCCAAGATITCAGATATICIGCTCTATIATATAAACTITATA
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               AAH13813 standard; cDNA; 2044
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The present invention describes primer sets for synthesising 5602 full-
length cDNAs defined in the specification. Where a primer set comprises:

[a) an oligo-dr primer and an oligonucleotide comprises one of the 5602
complementary strand of a polynucleotide which comprises one of the 5602
nucleotide sequences defined in the specification, where the
coligonucleotide comprises at least 15 nucleotides; or (b) a combination
of an oligonucleotide comprises a sequence complementary to the
complementary strand of a polynucleotide which comprises a 5'-end
complementary strand of a polynucleotide which comprises a 5'-end
complementary strand of a polynucleotide which comprises a 5'-end
complementary strand of a polynucleotide which comprises a 5'-end
complementary strand of a polynucleotide which comprises a 5'-end
complementary strand of a polynucleotide of sequence complementary to a
complementary strand of a polynucleotide of sequence of sequence of a polynucleotide of comprises at least 15 nucleotides and the combination of
coligonucleotide comprises at least 15 nucleotides and the combination of
coliforucleotide which comprises at least 15 nucleotides on those defined in the
coliforucleotide which comprises at least 15 nucleotides where the
coliforucleotide which set useful for synthesising polynucleotides,
coperitication and/or diagnosis of the abnormality of the proteins encoded by
the full-length cDNAs. The primers are also useful for the
detection and/or diagnosis of the abnormality of the proteins encoded by
the full-length cDNAs. The primers and allow obtaining of the full-length
connas and any specialised methods. AAH03166 to AAH13628 and
AAH13633 to AAH1872 represent human cDNA sequences; AAB93446 to AAB95893
crepresent human amino acid sequences, and AAH13629 to AAH13632 represent
complementary strands and the resemplification of the
complementary strands and the sequences and AAH13629 to AAH13632 represent
                                                                                                                                                                                                             Primer sets for synthesizing polynucleotides, particularly the 5602 full-length cDNAs defined in the specification, and for the detection and/or diagnosis of the abnormality of the proteins encoded by the full-length cDNAs.
                                                                                 <u>ب</u>
                                                                                    Yamamoto
                                                                                                                                                                                                                                                                                                                                                                                      Claim 8; SEQ ID NO 10767; 2537pp + Sequence Listing; English.
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                                                                                 Saito K,
Otsuki
                                                                                 Hayashi K, S
A, Nagai K,
                                                                                 hikawa T,
Wakamatsu
                                                                                 Nishikawa T, Wakamat
                        (HELI-) HELIX RES INST
                                                                                                                                                                        WPI; 2001-318749/34.
                                                                                                                   Sugiyama
                                                                                    Isogai T,
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                                                                                                                Ishii S,
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126 132 372 426 432 486 186 192 246 252 GTTAACTGCTAACTTTTCTGTAACCAACCTGAGTCAAACAGATGAAATCCCTGTATTGAC 306 312 366 CGCGGTTGCACGTCGGCCCCAGCCCTGAGGAGCCGGATGTGGAAACTGCTGCCCCCC CGCGGGCCCCGCAGGAGAACCATACAGACTTTTGACTGGCGTTGAGTACGTTGTTGG TTCCCGAACTTTGAAGTCGGGGATGGTATTACTTTTGGAGTGTTTGGAAGTAAATTCAG CGCGGGCCCGGCAGGAGGAGCATACAGACTTTTGACTGGCGTTGAGTACGTTGTTGG <u>AATAGAGTATGAGCCTTTGGTTGCATGCTCTTCTTGTTTTAGATGTCTCTGGGAAAACTGC</u> AAGGAAAAACTGTGCCATTCTAATTGAAAATGATCAGTCGATCAGCCGAAATCATGCTGT AAGGAAAAACTGTGCCATTCTAATTGAAATGATCAGTCGATCAGCCGAAATCATGCTGT GTTAACTGCTAACTTTTCTGTAACCAACCTGAGTCAAACAGATGAAATCCCTGTATTGAC CGCGGTTGCACGTCGGCCCCAGCCCTGAGGCCGGACCGATGTGGAAACTGCTGCCCGC Gaps DB 4; Length 2044; 1; Indels 119; 41.7%; Score 1836.4; 94.3%; Pred. No. 0; ive 0; Mismatches Matches 1976; Conservative Local Similarity 67 187 247 307 367 13 73 127 133 193 253 313 373 427 Query Match 원 ઠે a ò 셤 ઠે d ઠે 셤 8 유 ઠ à 움 433 TITAAATCAAGCTATATTGCAACTTGGAGGATTTACTGTAAACAATTGGACAGAAGAATG 492

99JP-00300253. 2000JP-00118776. 2000JP-00183767.

27-AUG-1999; 11-JAN-2000; 02-MAY-2000;

29-JUL-1999;

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28-JUL-2000; 2000EP-00116126

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                                                                                      The present invention describes primer sets for synthesising 5602 full-
length cDNAs defined in the specification. Where a primer set comprises:

(a) an oligo-dr primer and an oligonucleotide comprisentary othe
complementary strand of a polymucleotide which comprises one of the 5602
nucleotide sequences defined in the specification, where the
coligonucleotide comprises at least 15 nucleotides, or (b) a combination
of an oligonucleotide comprising a sequence complementary to the
complementary strand of a polymucleotide which comprises a 5'-end
sequence and an oligonucleotide comprising a sequence complementary to a
polymucleotide which comprises at least 15 nucleotides and the combination of
the 5'-end sequence/3'-end sequence; where the
oligonucleotide comprises at least 15 nucleotides and the combination of
the 5'-end sequence/3'-end sequence is selected from those defined in the
specification. The primer sets can be used in antisense therapy and in
gene therapy. The primer sets can be used in antisense therapy and in
gene therapy. The primers are useful for synthesising polymucleotides,
centection and/or diagnosis of the abnormality of the proteins encoded by
the full-length cDNAs. The primers are also useful for the
clust and primer are useful for synthesising polymucleotides,
che full-length cDNAs. The primers allow obtaining of the full-length
conna assily without any specialised methods. AAH03166 to AAH13628 and
AAH13633 to AAH13632 represent human amino acid sequences; and AAH13629 to AAH3632 represent
coligonucleotides, all of which are used in the exemplification of the
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defined in the specification, and for the detection and/or
the abnormality of the proteins encoded by the full-length
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                                                              2537pp + Sequence Listing; English.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                       Sequence 752 BP; 212 A; 158 C; 177 G; 202 T; 0 U; 3 Other;
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ilarity 99.4%; Pred. No. 1.1e-131;
Conservative 0; Mismatchhom
                                                              SEQ ID NO 495;
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es 682; Conserv
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               diagnosis of cDNAs.
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GACCTATTCCTGAAGCAGAAATTGGATTGGCGGTGATTTTCATGACTACAAAGAATTACT 1019
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 The invention relates to a method for the detection of cancer in which a change in the expression of 1 or more of 264 specified cancer associated genes, ABZ71694-ABZ71957, or of sequences at least 80% homologous to them in the specimen tissue as compared to normal tissue is observed. The genes are used in detection, diagnosis and treatment of cancer, especially of stomach cancer. The present sequence is that of a cancer associated polynucleotide of the invention
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        121 GACCTATTCCTGAAGCAGAAATTGGATTGGCGGTGATTTTCATGACTACAAAGAATTACT 180
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                                               671
                                                                                  Measuring changes in expression of 264 cancer associated genes for detection of stomach cancer and screening of potential anticancer agents.
607 TGGACGTCCAATTGTAAAGCCAGAATATTTTACTGAATTCCTGAAAGCAGTTCAGTCCAA
                                                  GAAGCAGCCTCCACAAATTGAAAGTTTTTACCCACCTCTTGATGAACCATCTATTGG-AA
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          Human; cancer; stomach cancer; cytostatic; gene;
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Pred. No. 3.2e-109;
0; Mismatches 11;
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                                                                                                                                                     GTAAAAATGTTGATCTGTCAGGACGG
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          Human cancer-associated gene
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21-SEP-2001; 2001JP-00290193.
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Best Local Similarity 97.6%;
Matches 604; Conservative
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Mori M;
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GIGATCCTCAGGGCCATCCCAGTACAGGATTAAAGACAACAACTCCAGGACCAAGCCTTT
                         CACAAGGCGTGTCAGTTGATGAAAACTAATGCCAAGCGCCCCAGTGAACACTACAACAT
                                     CACAAGGCGTGTCAGTTGAAAAAACTAATGCCAAGCGCCCCAGTGAACACTAACAACAT
                                                           ACGTAGCTGACACAGAATCAGAGCAAGCAGATACATGGGATTTTGAGTGAAAGGCCCAAAAG
                                                                             <u>accinecteacacacaarcagaccaaccagaracarceacartricacrcaaaacccaaaac</u>
                                                                                              AAATCAAAGTCTCCAAAATGGAACAAAATTCAGAATGCTTTCACAGAGGCGCCCACTG
                                                                                                        361 AAATCAAAAGTCTCCAAAAAGGAACAAAAATTCAGAATGCTTTTCACAAGACGGCCCACTG
                                                                                                                                                                   AAGATAGGGCTTNTTCAGCAGCAGCAACTTCCATCAGAACTACTTTCAGNCCG
                                                                                                                                TAAAGGAGTCCTGCAAAACAAGCTCTAATAATAATAGTATGGTATCAAATACTTTGGCTA
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          Sequences AAA77722-A78199 represent 478 cDNAs encoding proteins or
                                                                                                                                                                                                                                                                                                                                                                        colon tumour polypeptide; tumour antigen; cancer; vaccine;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      used to inhibit the development of for diagnosing and monitoring the
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                                                                                                                                                                                                                                                                                                                                                      CDNA encoding human colon tumour polypeptide, SEQ ID NO:343
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                                                                                                                                                                                                                                                                                                                                                                                  immunotherapy; diagnosis; progression; ss.
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                                                                                                                                                                                                                                       TCTACCAAAAAAAGGGAAA 1454
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99US-00347496.
99US-00401064.
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02-JUL-1999;
22-SEP-1999;
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Invention also specifically discloses 8 human colon tumours proteins invention also specifically discloses 8 human colon tumour proteins antigen presenting cells (APCS, preferably dendritic cells) expressing antigen presenting cells (APCS, preferably dendritic cells) expressing such polypeptides may be used in vaccines that target tumour cells, caper that target tumour cells, caper that target tumour cells, caper to cancer. T-cells specific for the polypeptide expressed by the APC are used to remove tumour cells from bological samples, especially blood or cancer. T-cells specific for the polypeptide expressed by the APC are used to remove tumour cells from bological samples, especially blood or caper then be used to inhibit cancer development. CD4+ and/or CD8+ T-cells from a patient may be incubated with a polypeptide or nucleic acid of the invention, or an APC expressing such a polypeptide, and then administered back to the patient to inhibit cancer development. Conclude endinistered back to the patient to inhibit cancer development. Conclude endinistered back to the patient to inhibit cancer development. The prospection of the invention, and therefore to determine whether cancer cancer cancer or progression of a cancer by repeating the processes time intervals, and comparing the current result to previous results. The present sequence represents a cDNA encoding a human colon tumour polypeptide
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              585 ACAGGCATTGAGCCAGTTAGATTATTGAAATATTATAGAGAGTTATGAACACTTAAATT
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of proteins which are associated with human colon tumours.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   Sequence 646 BP; 239 A; 124 C; 95 G; 188 T; 0 U; 0 Other;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Score 552.2; DB 3;
Pred. No. 1.4e-106;
4; Mismatches 32;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Query Match 12.5%;
Best Local Similarity 93.0%;
Matches 609; Conservative
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the polymucleotides (II) that encode them. (I) have cytostatic activity.

(I) and (II) can be used in gene therapy and vaccine production. (I) and (II) can be used in gene therapy and vaccine production. (I) and (II) can be used in gene therapy and vaccine production. (I) and (II) may be used in the prevention, diagnosis and treatment of diseases associated with inappropriate colon tumour associated protein (TCAP) cypression, such as colonic cancer. For example, (I) and (II) may be used to to treat disorders associated with decreased expression by rectifying mutations or deletions in a patient's genome that affect the activity of TCAPs by expressing inactive proteins or to supplement the patients own production of them. Additionally, (II) may be used to produce the TCAP proteins, by inserting the nucleic acids into a host cell culturing the proteins, by inserting the nucleic acids into a host cell culturing the cell to express the protein. (II) and its complementary sequences may also be used as DNA probes in diagnostic polymerase chain reaction (PCR) and hybridisation assays to detect and quantitate the presence of similar conclusion and activity. Anti-(I) and its conditions in the production of antibodies against TCAP expression and activity. The anti-(I) antibodies may also be used as diagnostic agents for detecting the cypression and activity. Anti-(I) antibodies and antagonists may also be used as diagnostic acids immunosorbant assay presence of TCAPs. All and AAL2841 and AAL2841 to AAAL2821 represent
                                                                                                                                                                                          Human; immunotherapy; diagnosis; colon cancer; colon tumour; immunogenic; gene therapy; vaccine; colonic cancer; ss.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        present invention describes colon tumour associated proteins (I) and
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 nucleotide and amino acid sequences given in the exemplification of the
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          Colon tumor associated proteins and nucleic acids useful for the prevention, diagnosis and treatment of colonic cancer.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Stolk JA;
                                                                                                                                                      Colon tumour related determined cDNA sequence for clone 25908
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      Sequence 646 BP; 239 A; 124 C; 95 G; 188 T; 0 U; 0 Other;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   Meagher MJ,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          Score 552.2; DB 4;
Pred. No. 1.4e-106;
4; Mismatches 32;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Benson DR,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    Claim 25; Page 228; 472pp; English.
RESULT 9
AA128794/c
ID AA128794 standard; cDNA; 646 BP.
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72000US-0048031.
2000US-00504629.
2000US-00519444.
2000US-0057251.
2000US-00674281.
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93.0%;
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                                                                                                                 (first entry)
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Best Local Similarity
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   Lodes MJ,
                                                                                                                                                                                                                                                                                     WO200149716-A2
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06-MAR-2000;
19-MAY-2000;
29-JUN-2000;
                                                                                                                                                                                                                                                    Homo sapiens.
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King G
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3718
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                                                                                                                                                                                                                    465 ATATTAGTTGAAAATGGAGTCATTTGAGTCTCTTAATAGCCATGTATCATAATTACCA 406
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King GE, Wang A, Clapper JD, Skeiky YAW;
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3659 AGTGAAGCTGGTGGAACATATGGTCTCCATTTTACAGTTAAGGAATATAATGGACAGATT
                                                                                                                                                                                                                                                                                                                                             AATATTGTTYTCTGTCATGCCCACAATCCCTTTCTAAGGAAGACTGCCCTACTATAGCAG
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                                                                              585 ACAGGCATTGAGACCAGTTAGATTATTGAAATATTATAGAGAGTTATGAACTTTAAATT
                                                                                                                            3539 ATGATAGTGGTATGACATTGGATAGAACATGGGATACTTTAGAAGTAGAATTGACAGGGC
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                                                       3479 ACAGGCATTGAGACCAGTTAGATTATTGAAATATTATAGAGAGTTATGAACACTTAAATT
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03-AUG-2001; 2001US-00922217.
19-DEC-2001; 2001US-00025380.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       30-JAN-2003 (first entry)
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Wang T,
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                                                                                                                        diagnosis of colon cancer. Also described: (1) a method for detecting the presence of cancer in a patient; (2) a method for stimulating and/or expanding T cells specific for a tumour protein; (3) an isolated T cell population comprising T cells prepared by the method of (2); (4) a method for stimulating an immune response in a patient; (5) a method for treating cancer in a patient; and (6) a method for inhibiting the development of cancer in a patient. (1) have immunostimulate and can be used in vaccines. AB232646 to AB233725 and ABE55343 to ABP55391 represent human colon cancer/tumour related
                                                                                                                                                                                                                                                                                                                                                                         586
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              406
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        170 TGATG-----TITCATTITGGITTTAATTIGTATATCCCTGATAGCTATAATTGGGTCAT 116
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         99
                                                                                                                                                                                                                                                                                                                                                 3419 AAAAGCTTCTCAGCCTTCCTAGGGAACAGAAATTGGGTGAGCCAATCTGCAATTTCTACT
                                                                                                                                                                                                                                                                                                                                                                     AAAGCTTTCTCAGCCTTCCTAGGGAACAGAAATTGGGGAGCCCAATCTGCAATTTCTAACT
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                                                                                                                                                                                                                                                                                                                                                                                                                  ATGATAGTGGTATGACATTGGATAGAACATGGGATACTTTAGAAGTAGAATTGACAGGGC
                                                                                                                                                                                                                                                                                                                                                                                                                                                                  ATATTAGTTGATGAAATGGAGTCATTTGAGTCTYTTAATAGCCATGTATCATAATTACCA
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   ATATTAGTTGATGAAATGGAGTCATTTGAGTCTCTTAATAGCCATGTATCATAATTACCA
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       AATATTGTTYTCTGTCATGCCCACAATCCCTTTCTAAGGAAGACTGCCCTACTATAGCAG
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           AATATTGTTCTCTGTCATGCCCACAATCCCTTTCTAAGGAAGACTGCCCTACTATAGCAG
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    CACCTAGTCTGTGGCTTGACTGTTTTCTTTATGTCTTTTGATGAATAGAAGTTTT 4073
                                                                                                                                                                                                                                                                                                                         Gaps
                                                                                                               present invention describes compounds (I) for the immunotherapy
                                                  New polynucleotide, useful for the preparation of a composition for stimulating an immune response against, or treating, cancer.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 CACCTAGTCTGTGGCTTGACTGTTTTCTTTATGTCTTTTTGATGAATAGAAGTTTT 1
                                                                                                                                                                                                                                                                                                                        Indels 10;
                                                                                                                                                                                                                                            sequences used in the exemplification of the present invention
                                                                                                                                                                                                                                                                                             Length 646;
                                                                                                                                                                                                                                                                     Sequence 646 BP; 239 A; 124 C; 95 G; 188 T; 0 U; 0 Other;
                                                                                                                                                                                                                                                                                            Query Match 12.5%; Score 552.2; DB 8; Best Local Similarity 93.0%; Pred. No. 1.4e-106; Matches 609; Conservative 4; Mismatches 32;
                                                                                         Example 1; Page 231; 537pp; English
  Carter
 Vedvick TS,
                          WPI; 2003-067548/06.
GR,
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expressed in cancer tissues. ABB7893 to ABB79004 represent proteins expressed in cancer tissues. ABB7893 to ABB79004 represent proteins encoded by the ABG60776 to ABG60787 uncleic acid sequences. (1) can be used in antisense therapy. An antibody immunoreactive with a polypeptide encoded by (1) is useful for detecting cancer in a patient sample, and for detecting the presence or absence of a polynucleotide encoded by a nucleic acid which hybridises to (1) in a cell. A probe/primer derived from (1) can be used for determining the presence of a nucleic acid which hybridises to (1), and for determining the phenotype of cells in a sample of colon cancer in a cell or tissue type, for determining the presence or state of other type of cancer, in antisense therapy, to generate macroarrays on a solid surface, to identify a chromosome on which the corresponding gene resides, and in tissue profiling, forensics, genetic analysis, mapping and diagnostic applications. (1) can be used to raise antibodies, and to screen for peptide analogues and antagonists
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      601
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           New isolated nucleic acid that is differentially expressed in cancer tissues useful for determining the presence of colon cancer in a cell or tissue type, and in antisense therapy.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      GCACTCATTIGIGGACGTCCAATIGIAAAGCCAGAATATTTACTGAATTCCTGAAAGCA
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  GCACTCATTTGTGGACGTCCAATTGTAAAGCCAGAATATTTTACTGAATTCCTGAAAGCA
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        CAGAAGAATGCACTCA-CCTTGTCATGGTATCAGTGAAAGTTACCATTAAAACAATATGT
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           561 CAGAAGAATGCACTCNCCCTTGTCATGGTATCAGTGAAAGTTACCATTAAACCAATATGT
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    GTTCAGTCCAAGAAGCAGCCTCCACAAATTGAAAGTTTTTACCCACCTCTTGATGAAGCA
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         Molino GA;
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                                                                                                                                                                                                                                                     Human; colon cancer; cancer; tissue profiling; forensic; mapping; genetic analysis; diagnostic; antisense therapy; gene; ss.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         1;
                                                                                                                                                                                         Human colon cancer related nucleotide sequence SEQ ID NO:1463.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Sequence 561 BP; 151 A; 120 C; 102 G; 187 T; 0 U; 1 Other;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         Dwivedi P,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         Indels
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Pred. No. 4.8e-105;
0; Mismatches 3;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         Catino TJ,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         Carroll E,
   BP.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   Claim 1; Fig 1; 796pp; English
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               02-OCT-2001; 2001WO-US030732.
ABQ57768 standard; cDNA; 561
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              02-OCT-2000; 2000US-0237271P.
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Best Local Similarity 99.3%;
Matches 557; Conservative
                                                                                                                              (first entry)
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          Lewis ME;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         Burgess C, Astle JH,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      WPI; 2002-426115/45.
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                                                                                                                                                                                                                                                                                                                                                                                                                 WO200229086-A2.
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                                                                                                                                                                                                                                                                                                                                                     Homo sapiens.
                                                                                                                              02-AUG-2002
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                11-APR-2002
                                                                  ABQ57768;
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Gaps

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3610 TGAAATGGAGTCATTTGAGTCTYTTAATAGCCATGTATCATAATTACCAAGTGAAGCTGG 3669
                                                                                                                     3190 AGAGAATGAGAAATGGAACAGTGAGGAATGGAGGCCATATTTCCATGACTTCCCTTGTAA 3249
                                                                                                                                                                                                                                                                                                                             241 AGGIGGAACTCCAGCIGCAAGGGAGTTAGGGAAATGAAGGICTTTTTTAAAAGCTTCTC 300
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          Human; primer; detection; diagnosis; antisense therapy; gene therapy; ss.
                                                                                                                                      AGAGAATGAGAAATGGAAACAGTGAGGAATGGAGGCCATATTTCCATGACTTCCCTTGTAA
                                                                                                                                                                                                121 ACAGAAGCAACAGAAGGGACAAGAGGCTGGCCTCTACATCACTCTCAAATCTT
                                                                                                                                                                                                                                        GTGGAAGTGCATTCCTACCAGAACCAAATTAACTTACTTCCAAGTTCTGGCTTGC
                                                                                                                                                                                                                                                                      GreenAgrecarcracardocagaaccaaarraacrracarcracarrorgecrecardocage
                                                                                                                                                                                                                                                                                                                                                                                        301 AGCCTTCCTAGGGAACAGACATTGGGTGAGCCAATCTGCAATTTCTACTACAGGCATTGA
                                                                                                                                                                                                                                                                                                                                                                                                                                      361 GACCAGTTAGATTATTGAAATATTATAGAGAGTTATGAACACTTAAATTATAGAGGGT
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              TGAAATGGAGTCATTTGAGTCTCTTAATAGCCATGTATCATAATTACCAAGTGAAGCTGG
                                                                              250 ACAGAAGCAACAGAAGGGACAAGAGGCTGGCCTCTACATCACTCTCACCTTCCAAATCTT
                                                                                                                                                                                                                                                                                                   AGGIGGAACTCCAGCTGCAAGGGAGTTAGGGAAATGAAGGTCTTTTTAAAAGCTTCTC
                                                                                                                                                                                                                                                                                                                                                              3430 AGCCTTCCTAGGGAACAGAAATTGGGTGAGCCAATCTGCAATTTCTACTACAGGCATTGA
                                                                                                                                                                                                                                                                                                                                                                                                                     GACCAGTTAGATTATTGAAATATATAGAGAGTTATGAACACTTAAAATTATGATAGTGGT
                                                                                                                                                                                                                                                                                                                                                                                                                                                                              ATGACATTGGATAGAACATGGGATACTTTAGAAGTAGAATTGACAGGGCATATTAGTTGA
                                                             ACCTGTGATCCAGCAAGAAGGGAGTTCCAGTCAAGAGTCACTACAACTGATTAGTTGTTT
                                ö
 Length 544;
                                Indels
Score 538.6; DB 6;
Pred. No. 9.7e-104;
2; Mismatches 3;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Human cDNA clone (3'-primer) SEQ ID NO:6023
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         Bb.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      29-JUL-1999; 99JP-00248036.
27-ANG-1999; 99JP-00300253.
11-JAN-2000; 2000JP-00118776.
02-MAY-2000; 2000JP-00183767.
09-JUN-2000; 2000JP-002418999.
   12.2%;
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                              Matches 539; Conservative
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   Query Match
Best Local Similarity
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AAH09188/
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                                                                                                                                                                                                                                        CCTATTCCTGAAGCAGAAATTGGATTGGCGGTGATTTTCATGACTACAAAGAATTACTGT 1021
                                                                                                                      901
                                                                                                                                                    142
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               ABL36412 to ABL38645 represent human colon tumour antigen cDNA clones (I) which were isolated from human colon tumour and colon metastatic tumour cDNA libraries. (I) have cytostatic activity and can be used in vaccine production. (I) can be used for stimulating and/or expanding T cells specific for a tumour protein on contact with the T cells. They are also useful for inhibiting the development of cancer in a patient. (I) can be used as probes or primers for nucleic acid hybridisation, for preparing can be used in the diagnosis of a colon tumour
                                                                                                                                                                                 GACTGTCAGAAGAAATGGATTCAGTCAATAATGGATATGCTCCAAAGGCAAGGTCTTAGA 961
                                                                                                                                                                                                           82
                                                                                                                                                                                                                                                                      22
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          Novel isolated colon tumor polynucleotide differentially expressed in colon tumor or colon metastatic tumor and polypeptides encoded by them, useful for inhibiting development of cancer in patient.
                    321 GGGAAAACATTTATATATTTTTGAATGCCAAACAGCATAAGAAATTGAGTTCCGCAGTTGTC
                                                                                                                                                                                                  GACTGTCAGAAGAAATGGATTCAGTCAATAATGGATATGGATCTAGAAGGCAAGGTCTTAGA
                                                                                                                        TTGGCTCCGGGAACGTGTTGTTGATACAGGAATAACAAACTCACAGACCTTAATTCCT
                                                                                                                                       201 TTGGCTCCGGGAACGTGTGTTGTTGATACAGGAATAACAAACTCACAGACCTTAATTCCT
                                                                                                                                                                                                                                                           261 TTTGGAGGTGGGGAAGCTAGTTGATAACAGAAGAAGAAGAAGAAGAAGAAGAATTTCTTT
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             Human; colon cancer; colon tumour antigen; cytostatic; vaccine; colon tumour metastatic antigen; diagnosis; gene; ss.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Sequence 544 BP; 173 A; 94 C; 125 G; 150 T; 0 U; 2 Other;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               Human colon tumour antigen polynucleotide SEQ ID NO:1737.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     Claim 1; SEQ ID NO 1737; 105pp; English
                                                                                                                                                                                                                                                                                                   GATCCTCAGGGCCATCCCAGT 1042
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                                                                                                                                                                                                                                                                                                                       GATCCTCAGGGCCATCCCAGT
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20-FEB-2001; 2001US-0270216P.
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line present interaction describes primer acts synthemestations. The present interaction describes primer acts solutions and an oligonucleotide complementary to the complementary strand of a polyuncleotide which comprises one of the 5602 nucleotide sequences defined in the specification, where the oligonucleotide sequences defined in the specification, where the oligonucleotide comprises at least 15 nucleotides, or (b) a combination of an oligonucleotide comprising a sequence complementary to the complementary strand of a polyuncleotide which comprises a 5'-end sequence and an oligonucleotide comprising a sequence complementary to a polynucleotide comprises a 1'-end sequence, where the polynucleotide comprises a 1'-end sequence, where the combination of the 5'-end sequence/3'-end sequence is selected from those defined in the specification. The primer sets can be used in antisense therapy and in gene therapy. The primer set useful for synthesising polynucleotides, compared the full-length cDNAs. The primers are useful for synthesising polynucleotides and detection and/or diagnosis of the abnormality of the proteins encoded by the full-length cDNAs. The primers allow obtaining of the full-length CDNAs and seasily without any specialised methods. AAH03163 to AAH3633 represent human amino acid sequences; and AAH3632 to AAH3632 represent contaction of the contac particularly the 5602 full-Primer sets for synthesizing polynucleotides, particularly the 5602 full length cDNAs defined in the specification, and for the detection and/or diagnosis of the abnormality of the proteins encoded by the full-length present invention describes primer sets for synthesising 5602 full-Yamamoto J; Claim 3; SEQ ID NO 6023; 2537pp + Sequence Listing; English. Saito K, Ya Otsuki T; Hayashi K, S A, Nagai K, Nishikawa T, T, Wakamatsu WPI; 2001-318749/34. Isogai T, N , Sugiyama 1 present invention Ota T, I Ishii S, 

Sequence 543 BP; 128 A; 111 C; 83 G; 219 T; 0 U; 2 Other;

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1; 1683 1743 1803 1863 1923 1983 2043 1564 GGAGCAGCATCTATCTGAGAATGAGCCTGTGGACACAAACTCAGACAATAACTTATTTAC 1623 485 425 365 245 305 185 184 TGAACTCAAGGAAGACTCACTATGGTCAGCTAAAGAAATATCTAACAATGGCAAACTTCA 125 65 GNAGCAGCTTCTATCTGAGAATGAGCCTG-GGACCCAAACTCAGCCAATAACTTATTTCC **AGATACAGATTTAAAATCTATTGTGAAAAATTCTGCCAGTAAATCTCATGCTGCAGAAAA** AGATCCAGATTTAAAATCTATTGTGAAAAATTCTGCCAGTAAATCTCATGCTGCAGAAAA GCTAAGATCAAATAAAAAAGGGAAATGGATGATGTGGCCATAGAAGATGAAGTATTGGA GCTAAGATCAAATNAAAAAAGGGAAATGGATGTGGCCATAGAAGGTATATGGA ACAGTTATTCAAGGACACAAAACCAGAGTTAGAAATTGATGTGAAAGTTCAAAAACAGGA GGAAGATGTCAATGTTAGAAAAAGGCCAAGGATGGATATAGAAACAAATGACACTTTCAG TGATGAAGCAGTACCAGAAAGTAGCAAAATATCTCAAGAAATGAAATTGGGAAGAAACG TGATGAAGCAGTACCAGAAAGTAGCAAAATATCTCCAAGAAAATGAAATTGGGAAGAAACG TGAACTCAAGGAAGACTCACTATGGTCAGCTAAAGAAATATCTAACAATGACAAACTTCA .984 GGATGATAGTGAGATGCTTCCAAAAAGCTGTTATTGACTGAATTTAGATCACTGGTGAT 124 GGATGATAGTGAGTGCTTCCAAAAAGCTGTTATTGACTGAATTTAGATCACTGGTGAT Gaps 1; Length 543; Indels DB 4; .. 6 Score 520.4; DB 4; Pred. No. 6.8e-100; 0; Mismatches 11.8%; 98.3%; Conservative Local Similarity Best Local Simi Matches 535; 543 1624 1684 424 1744 364 1804 304 244 484 1864 1924 Query Match 셤 셤 ð 셤 g 윱 g 셤

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The invention relates to a new isolated polymucleotide (a Human kidney tumour specific CDNA) comprising any one of the 1855 sequences identified in the specification (or their complements, degenerate variants, sequences consisting of at least 20 contiguous residues them, sequences that hybridise to them under highly stringent conditions or sequences that hybridise to them under highly stringent conditions or sequences consisting at least 75 or 90% sequence identity to the 1855 sequences. Also included are detecting/determining the presence of cancer in a patient, an isolated polypeptide encoded by one of the 1855 sequences, an expression vector comprising the polymucleotide operably linked to an expression control sequence, a host cell transformed/transfected with the vector, an isolated antibody (or its antigen-binding fragment) that specifically binds to the protein, a fusion protein comprising at least one the protein, an isolated T-cell population comprising at least one the protein. Comprising a first component (comprising the T-cells, a composition as second component (comprising one of the polymulation as second component (comprising and a diagnostic kit presenting cell that expresses the polymucleotide) and a diagnostic kit comprising at least one of the oligonucleotides, or at least one antibody are presenting cell that expresses the polymucleotide) and a diagnostic kit comprising at least one of the oligonucleotides, or at least one of the oligonucleotides, or at least one of the presenting cancer by stimulating and/or expanding T-cells specific for a tumour comprising a response in a patient. The present compression and attending immune response in a patient. The present

Claim 1; SEQ ID NO 532; 78pp; English.

response in a patient.

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  TAAAAACTCTACTTCCAGAAATCCGTCTGGCATAAATGATGATTATGGTCAACTAAAAAA
                                              64 TAAAAACTCTACTTCCAGAAATCCGTCTGGCATAAATGATGATTATGGTCAACTAAAAA
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21-DEC-2001; 2001US-0343340P.
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CTTCCCTTGTAAACAGAAGCAACAGAAGGGACAAGAGGCTGGCCTCTACATCACTCTCAC 3297
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               88; sequencing by hybridisation; SBH; expressed sequence tag; BST;
mapping; biodiversity; genetic disorder.
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sequence data for this patent did not form part of the printed specification, but was obtained in electronic format directly from USPTO at seqdata.uspto.gov/sequence.html?DocID=20030109434.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         AGAAATTCTTTATACATTCTAGATGCAAGTCTCTTGYCGGATATACGTATTGAGATA 4015
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                                                                                                   Gaps
                                                                                                 10;
                                                                         Length 587;
                                                  Sequence 587 BP; 177 A; 86 C; 114 G; 210 T; 0 U; 0 Other;
                                                                                                24; Indels
                                                                        Query Match 11.5%; Score 507; DB 7;
Best Local Similarity 93.6%; Pred. No. 4.7e-97;
Matches 559; Conservative 4; Mismatches 24;
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genome
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The invention relates to an isolated polynucleotide comprising any one of 38043 CDNA sequences, appearing as ACH12789-ACH50831, whose sequence was determined by the technique of SBH (sequencing by hybridisation). Also included is a purified polypeptide comprising a sequence corresponding to a reading frame of the novel polynucleotide. The nucleic acid sequences are useful in diagnostics as expressed sequence tags (BST) for identifying expressed games or for physical mapping of the human genome, in forensics, in assessing biodiversities, or in identifying mutations responsible for genetic disorders and other traits. The nucleotide sequences are also useful as hybridisation probes, as oligomers for PCR, for chromosome and gene mapping, in the recombinant production of protein, or in generating antisense DNA or RNA. The purified polypeptide is useful for generating antibodies specific for it. The present sequence is one of the 38043 isolated cDNA/BST sequences. Note: The sequence data for the thin patent did not form part of the printed specification, but was obtained in electronic format directly from USPFO at sequence.html?DocID=20030073623
                                                                                                                                                                                                                                                                                                                                                                 New polynucleotide sequences obtained from various cDNA libraries, useful as hybridization probes, as oligomers for PCR, for chromosome and gene mapping, in the recombinant production of protein, or in generating
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Pred. No. 1.2e-75;
1; Mismatches 10;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        Claim 1; SEQ ID NO 13834; 44pp; English
                                                                                                                                                                                                                        Stache-Crain B,
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Matches 411; Conservative
DRMANAC R T.
LABAT I.
STACHE-CRAIN B.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   antisense DNA or RNA.
                                                                                                          DICKSON M C.
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Search completed: August 30, 2005, 06:01:37 Job time : 2165 secs

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Sequence la Sequen

16201,

16016,

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14957, 50509,

Run on:

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GTCGGGGGATGGTATTACTTTTGGAGTGTTTGGAAGTAAATTCAGAATAGAGTATGAGCC 387
                                                                                                                                                                                                                                                                                                                             Pacent No. 6458534
GENERAL INFORMATION:
APPLICANT: CONCANNON Et al.
TITLE OF INVENTION: AGENE ASSOCIATED WITH NIJMEGEN BREAKAGE
TITLE OF INVENTION: SYNDROME, ITS GENE PRODUCT AND METHODS FOR THEIR USE
FILE REFERENCE: 9924-0003-228
CURRENT APPLICATION NUMBER: US/09/300,008B
CURRENT FILING DATE: 1999-04-27
PRIOR FILING DATE: 1998-04-27
PRIOR FILING DATE: 1998-04-27
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           TTCTGTAACCAACCTGAGTCAAACAGATGAAATCCCTGTATTGACATTAAAAGATAATTC
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US-09-949-016-12791

US-09-949-016-12793

US-09-949-016-12793

US-09-949-016-16014

US-09-949-016-16015

US-09-949-016-16015

US-09-949-016-16202

US-09-949-016-16203

US-09-949-016-16203

US-09-949-016-16203

US-09-949-016-1774

US-09-949-016-1776

US-09-949-016-1776

US-09-949-016-1776

US-09-949-016-1776

US-09-949-016-1776

US-09-949-016-17808

US-09-949-016-17808

US-09-949-016-17808

US-09-949-016-17808
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Score 4108.4;
Pred. No. 0;
7; Mismatches
                                                                                                                                                                                                                                                                    ALIGNMENTS
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SOFTWARE: PastSEQ for Windows Version 3.0
SEQ ID NO 1
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Best Local Similarity 97.1%;
Matches 4272; Conservative
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LOCATION: (26)...(2287)
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  US-09-300-008B-1
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          LENGTH: 4386
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Sequence 343, App
Sequence 14033, A
Sequence 12777, A
Sequence 12777, A
Sequence 1255, A
Sequence 12055, A
Sequence 13055, A
Sequence 1379, A
Sequence 12449, A
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    22, Appl
16676, A
152075,
152148,
                                                                                        August 30, 2005, 05:25:38 ; Search time 677 Seconds (without alignments) 10641.841 Million cell updates/sec
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Sequence 12396,
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1 ttcggcacgaggcgcggttg.....accgcggtggagctccagct
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Sequence
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(cgn2_6/ptodata/1/ina/5A_COMB.seq:*
(cgn2_6/ptodata/1/ina/5B_COMB.seq:*
(cgn2_6/ptodata/1/ina/6A_COMB.seq:*
(cgn2_6/ptodata/1/ina/6B_COMB.seq:*
(cgn2_6/ptodata/1/ina/PCTUS_COMB.seq:*
               5.1.6
Compugen Ltd.
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US-09-949-016-159526
US-09-949-016-159599
US-09-949-016-12396
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US-09-401-064-343
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US-09-533-080A-425
US-09-949-016-12777
US-09-949-016-12777
US-09-949-016-1379
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US-09-949-016-12784
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US-09-949-016-12218
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                                                                                                                                                                                                                                        1202784 seqs, 818138359 residues
               version 9
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Maximum Match 100%
Listing first 45 summaries
                                                                 nucleic search, using sw model
                                                                                                                                                                                                IDENTITY NUC
Gapop 10.0 , Gapext 1.0
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Copyright (c) 1993
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seq length: 2000000000
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552.2
84
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75.4
669.6
68.2
67.8
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Maximum DB 8
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Database

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us-09-837-138-1.rni

1468 TOMGAMPTICTICATIOCAMPTORATIONALINAMACTICTICATITICATION   111
8 4 8 4 8 6 8 6 8 6 8 6 8 6 8 6 8 6 8 6
10   STOCGGGGATGCTTTTTCGAGGTGTTTGGAAGTAATTCGAGATAAGATTTTTTTT

	; CURRENT FILING DATE: 1999-09-22
1	3549 TATGACATTGGATAGAACATGGGATACTTTAGAAGTAGAATTGACAGGGCATATTAGTTG 3608

34;

Length 601; Indels

1.9%; Score 84; DB 4; Le 56.6%; Pred. No. 2.9e-11; live 3; Mismatches 153;

------AATTTAACCCCAAGATTTCAGATATT

us-09-837-138-1.rni

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3945 TATAATTGGGTCATAGAAATTCTTTATACATTCTAGATGCAAGTCTCTTGYCGGATATAC 4004
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                                                                                                                                                                                                                                                                 120 ACTGTGA-ATTTTCTCCTTTGTCTTGTCTATTTATTCCTTAATGGTATCTTTTGAAAGC 178
                                                                                                                                                                                                                                                                                                                                                                                                                      179 AGAAGTTATAAATACTGATAGTGTCCAATTTATCTTTTCATTTGTTTAGTGCTTTCTTG 238
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                                                                                                                                                                                                                                                                                                              4005 GTATTGAGATATTACACCTAGTCTGTGACTTTTTCTTTTATGTCTTTTGATGAAT
                                                                                                                                                                                                                                                                                                                                                                                      TATGTTGTGAATTATGGATCAGGGTTCTTTTTTTTCCCCCATACAAGTATCCAGTCATTGT
 PRIOR FILING DATE: 2000-09-08
NUMBER OF SEQ ID NOS: 207012
SOFTWARE: FastSEQ for Windows Version 4.0
SEQ ID NO 79057
LENGTH: 601
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                                                                                                                                                                                      Best Local Similarity 56.6
Matches 248; Conservative
                                                                                                               ORGANISM: Human
                                                                                                                               US-09-949-016-79057
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                                                                                                                                                                                                                                                                                                                                                                       ATGATAGTGGTATGACATTGGATAGAACATGGGATACTTTAGAAGTAGAATTGACAGGGC 3598
                                                                                                                                                                                                                                                                                                                                                                                                                                             ATATTAGFTGATGAAATGGAGTCATTTGAGTCTYTTAATAGCCATGTATCATAATTACCA 3658
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  AGTGAAGCTGGTGGAACATATGGTCTCCATTTTACAGTTAAGGAATATAATGGACAGATT 3718
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      AATATTGTTYTCTGTCATGCCCACAATCCCTTTCTAAGGAAGAAGACTGCCCTACTATAGCAG 3778
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     3839 ACAAATATTGGGTGTTGTCCAGTATTTTTCCCTTTTTAACCMTTCCCAATTCGGGTGTGT 3898
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           3899 AGGIGGATGTTTCCATTTGGGTTTTAATTTGTATATCCCTGATAGCTATAATTGGGTCAT 3958
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          345 ANTATTGTTCTCTCTCATGCCCACAATCCCTTTCTAAGGAAGACTGCCCTACTATAGCAG 286
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     226 ACAAATATTGG----TGTTGTCAGTATTTTTTCCTTTTTTAACCATTCCAATCGGTGTGTAG 171
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      26
                                                                                                                                                                                                                                                            645 AAAGCTTTCTCAGCCTTCCTAGGGAACAGAAATTGGGGAGCCCAATCTGCAATTTCTACT
                                                                                                                                                                                                                                                                                                                                                                                          525 ATGATAGTGGTATGACATTGGATAGAACATGGGATACTTTAGAAGTAGAATTGACAGGGC
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   115 AGAAATTCTTATACATTCTAGATGCAATTCTCTTGTGGGATATATGTGATGTATTTA
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                                                                                                                                                                                        Gaps
                                                                                                                                                                                      10;
                                                                                                                                                    Length 646;
                                                                                                                                                                                      Indels
                                                                                                                                                  DB 4;
                                                                                                                                                                                        32;
                                                                                                                                                  Score 552.2; DB 4;
Pred. No. 4.5e-131;
4; Mismatches 32;
NUMBER OF SEQ ID NOS: 371
SOFTWARRE: FastSEQ for Windows Version 3.0
SEQ ID NO 343
LENGTH: 646
                                                                                                                                                  Query Match
Best Local Similarity 93.0%;
Matches 609; Conservative
                                                                           TYPE: DNA
CORGANISM: Homo sapien
US-09-401-064-343
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US-09-949-016-79057
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us-uy-say-uso-livus;

patent No. 681239

GENERAL INFORMATION:

APPLICANT: VENTER, J. Craig et al.

APPLICANT: VENTER, J. Craig et al.

TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED

TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF

FILE REPRENCE: CL001307

CURRENT APPLICATION NUMBER: US/09/949,016

CURRENT PILING DATE: 2000-04-14

PRIOR PAPLICATION NUMBER: 60/231,768

PRIOR PLILNG DATE: 2000-10-20

PRIOR PLILNG DATE: 2000-10-03

PRIOR FILING DATE: 2000-09-08

NUMBER OF SEQ ID NOS: 2000-10-03

PRIOR FILING DATE: 2000-09-08

NUMBER OF SEQ ID NOS: 207012

SOFTWARE: FastSEQ for Windows Version 4.0

LENGTH: 784019
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LOCATION: (1)...(784019)
OTHER INFORMATION: n = A,T,C or
RESULT 4
US-09-949-016-14033
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 TYPE: DNA
ORGANISM: Human
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Sequence 79057, Application US/09949016
Sequence 79057, Application US/09949016
Batent No. 6812339
Batent No. 6812339
TITLE OF INVENTER, J. Craig et al.
TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF FILE REFERENCE: CLO01307
CURRENT APPLICATION NUMBER: US/09/949,016
CURRENT APPLICATION NUMBER: 60/241,755
PRIOR FILING DATE: 2000-10-20
PRIOR PELICATION NUMBER: 60/231,768
PRIOR PELICATION NUMBER: 60/231,768
PRIOR PELICATION NUMBER: 60/231,498
PRIOR APPLICATION NUMBER: 60/231,498

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TYPE: DNA ORGANISM: Homo sapiens
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Sequence 12777, Application US/09949016

Sequence 12777, Application US/09949016

Sequence 12777, Application US/09949016

Sequence 12777, Application US/09949016

GENERAL INFORMATION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED

TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF

TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF

FILE REPERENCE: 2000-04-14

CURRENT APPLICATION NUMBER: 60/231,768

PRIOR FILING DATE: 2000-10-03

PRIOR APPLICATION NUMBER: 60/231,498

PRIOR FILING DATE: 2000-09-08

NUMBER OF SEQ ID NOS: 207012

SEQ ID NO 12777

LENGTH: 828152
                                                                                                                                                              167506 TTCTAAGAACCTTTACCAATCACCAGGTCACAAAAATATTTTCCTATGTTTTCTTCTAGA 467565
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                                                                                                                                                                                                                                             167387 ACTGTGA-ATTTTCTCCTTTGTCTTGTCTATTTATTCCTTAATGGTATCTTTTGAAAAGC 467445
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                                                                                                                     3945 TATAATTGGGTCATAGAAATTCTTTATACATTCTAGATGCAAGTCTCTTGYCGGATATAC 4004
                                                                                                                                                                                                        4005 GTATTGAGATATTACACCTAGTCTGTGGCTTGACTGTTTTCTTTATGTCTTTTGATGAAT
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                                                                                 34; Gaps
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                                    Length 784019;
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56.6%; Pred. No. 3.7e-09;
tive 2; Mismatches 154; Indels 34;
                                    Score 82.8; DB 4; Length 70
Pred. No. 3.6e-09;
2; Mismatches 154; Indels
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OTHER INFORMATION: n = A,T,C or
                                      1.9%;
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Best Local Similarity 56.61
Matches 248; Conservative
                                                                               Conservative
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                                                           Best Local Similarity
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   ORGANISM: Human
US-09-949-016-14033
                                                                               Matches 248;
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3945 TATAATTGGGTCATAGAAATTCTTATACATTCTAGATGCAAGTCTCTTGYCGGATATAC 4004

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APPLICANT: JOAN, KNOLL
APPLICANT: JOAN, KNOLL
APPLICANT: GOOSAN, PETER
TITLE OF INVENTION: SINGLE COPY GENOMIC HYBRIDIZATION PROBES AND METHOD OF GENERAT
FILE REPERENCE: 30307
CURRENT APPLICATION NUMBER: US/09/573,080A
CURRENT FILING DATE: 2000-05-16
NUMBER OF SEQ ID NOS: 479
SOFTWARE: Patentin version 3.0
SEQ ID NO 425
463819 AGCACTATTTGTTCAAAGACTAATCCTCGCGCTATTCAATTTCCTTGGCATCTTTGTCAA 463878
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             463639 TTCTAAGAACCTTTACCAATCACCAGGTCACAAAAATATTTTCCTATGTTTTCTAGA 463698
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DATABASE ENTRY DATE:
DATABASE ENTRY DATE: 1996-01-26
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           55,
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... Sequence 425, Application US/09573080A
... Patent No. 6828097
... GENERAL INFORMATION:
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LOCATION: (1). (3285)
OTHER INFORMATION: 11me_orf2
NAME/KEY: misc_feature
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RESULT 8
US-09-949-016-14061/c
Sequence 14061, Application US/09949016
Patent No. 681239
GENERAL INFORMATION:
APPLICANT: VENTER, J. Craig et al.
APPLICANT: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
FILE REFERENCE: CL001307
                                                                                                                                                                                                                                                                                                                                                                                                                                                 1677 CAGAAAAGCTAAGATCAAATAAAAAAGGGAAATGGATGATGTGGCCATAGAAGATGAAG 1736
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                                                                                                                                                                                                                                                                                                                   Length 7218;
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1.8%; Score 79.8; DB 1; Length 7
Best Local Similarity 3.8%; Pred. No. 1.4e-09;
Matches 15; Conservative 244; Mismatches 136; Indels
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CURRENT FILING DATE: 2000-04-14
PRIOR APPLICATION NUMBER: 60/241,755
PRIOR PILING DATE: 2000-10-20
PRIOR PLILING DATE: 2000-10-30
PRIOR APPLICATION NUMBER: 60/231,768
PRIOR PILING DATE: 2000-10-03
PRIOR PRIOR DATE: 2000-09-08
NUMBER OF SEQ ID NOS: 207012
SOFTWARE: PREUSEQ for Windows Version 4.0
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NAME/KEX: misc_feature
LOCATION: (1)...(75216)
OTHER INFORMATION: n = A,T,C or G
                                          TELEX: 899149
INFORMATION FOR SEQ ID NO: 3
SEQUENCE CHARACTERISTICS:
LENGTH: 7218 base pairs
                         (703)683-4109
                                                                                                                                          TYPE: nucleic acid
STRANDEDNESS: single
                                                                                                                                                                                                                                     , CLONE: pTZgpt-F1s
US-08-232-463-14
                                                                                                                                                                                             TOPOLOGY: linear IMMEDIATE SOURCE:
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TELEPHONE:
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                           TELEFAX:
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                                3203 TITITAATIGGGITAITIGGCTTTTTAITGAGTIGTAAGAGTICTTTATATATATGTG 3144
                                                                                                                           3143 GATACAAGTCCTTTGTCAGATATATGWTTTGCAAATATTTTCTCCCAGTCTGTGGGCTTGT 3084
                                                                                                                                                                                               CTGTTTTCTTTA-----TGTCTTTTGATGAATAGAAGTTTTAAATTTTGACAAGGTCA 4090
                                                                                                                                                                                                                                                                                             4091 AATTIAT---------TITTITCTTTTGTTTGATATTTTTCTCTCCAAT 4131
                                                                                                                                                                                                                                                                                                                                                                                            4132 TTAACCCCAAGATTTCAGATAT-----TCTGCTCTATTATAAAACTTTATATTTTA 4184
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TITIAAITITGIAIATCCCTGAIAGCIAIAAITGGGICAIAGAAAITCTTIAIACAITCIA 3979
                                                                                                                                                                                                                                          3083 criricariricrigarigicririrgaagagcagaagririraaririgargaagrcc 3024
                                                                                                                                                                                                                                                                                                                        3023 AATTTATCAATTTTTCCTTTTGTTGCTTGTGCTTTTKGTGTCATATCTAAGAAATCWTTG
                                                                                                                                                                                                                                                                                                                                                                                                                        2963 CCTARCCCAAGGICACGAAGATTTTCTCCTATGTTTTCTTCTAARAGTTTTATAGTTTTA
                                                                                                  980 GATGCAAGTCTTTGYCGGATATACGTATTGAGATA--TTACACCTAGTCTGTGGCTTGA
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OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/232,463
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Sequence 14, Application US/08232463
Patent No. 5670367
GENERAL INFORMATION:
APPLICANT: SCHEIFLINGER, F.
APPLICANT: FALTORER, F. G.
TITLE OF INVENTION: RECOMBINANT FOWLPOX VIRUS
NUMBER OF SEQUENCES: 52
CORRESPONDENCE ADDRESS:
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        FILING DATE:
APPLICATION NUMBER: EP 91 114 300.6
FILING DATE: 26-AUG-1991
ATTORNEY AGENT INPORMATION:
NAME: BENT, Stephen A.
REGISTRATION NUMBER: 39,768
REFERENCE/DOCKET NUMBER: 30472/114 IMMU
TELECOMMUNICATION INPORMATION:
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          ADDRESSEE: Foley & Lardner
STREET: 1800 Diagonal Road, Suite 500
CITY: Alexandria
STATE: VA
COUWIRY: USA
ZIP: 22313-0299
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PRIOR APPLICATION DATA:
APPLICATION NUMBER: US/07/935,313
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           ZIP: 22313-022.
COMPUTER READABLE FORM:
TYPE: Floppy disk
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APPLICATE: VENTER, J. Craig et al.,
APPLICATE: VENTER, J. Craig et al.,
ITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
ITTLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF;
ITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF;
CURRENT APPLICATION NUMBER: US/09/949,016
CURRENT FILING DATE: 2000-04-14
PRIOR APPLICATION NUMBER: 60/231,768
PRIOR PILING DATE: 2000-10-03
PRIOR FILING DATE: 2000-10-03
PRIOR FILING DATE: 2000-09-08
NUMBER OF SEQ ID NOS: 207012
SEQ ID NO 17190
ILENGTH: 141455
126483 TAATGAGCAGAGGITCTCATITITTATGAGATGTÄÄTTÄÄTTAACTITCTTTAATGAT 126542
                                                                                                                                                                                                                                    126543 TATTAATTTCTGTGTTCTAAGAAAGCTTTGCTTACCCATAGTCAGGAAGGTTCACCTTTG 126602
                                                                                                                                                                                                                                                                                                                                           126603 TTTAGGCCTATAGTCCATCTCTTTTTTTTTTTCATGAATGGTGTAGTAGTAGAAATTGA 126662
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                                                                                                                                                                                4117 ITITITICICICCAATITAACCCCAAGATITICAGATATICIGCICTATIATAAACTITA 4176
                                                                              4177 TATTTTTATATTTGTGATCTACCTTGAATTGAT--ATGTATGTTGTGAATTATGGATCAG 4234
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 126423 GTTTTGCAAATACTTTTCCCCAGTCTGCATGCATGACAATTTGTTTTTTGTGAAGTGTTT
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55.1%; Pred. No. 1.1e-07;
ive 1; Mismatches 162; Indels
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              ; Sequence 17190, Application US/09949016; Patent No. 6812339; GENERAL INFORMATION:
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           ; LOCATION: (1)...(141455)
; OTHER INFORMATION: n = A,T,C or G
US-09-949-016-17190
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Matches 212, Conservative
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                             126423
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TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
FILE REPERENCE: CLOO1307
CURRENT APPLICATION NUMBER: US/09/949,016
CURRENT APPLICATION NUMBER: US/09/949,016
PRIOR APPLICATION NUMBER: 60/241,755
PRIOR APPLICATION NUMBER: 60/241,755
PRIOR PILING DATE: 2000-10-20
PRIOR PILING DATE: 2000-10-03
PRIOR PILING DATE: 2000-10-03
PRIOR PILING DATE: 2000-09-08
NUMBER OF SEQ ID NOS: 207012
SOFTWARE: FASESEQ for Windows Version 4.0
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                                                                                                                                                                                                                                                                                                                681
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                                                                                                                                                                                        796 TATTATTGAGTTGTAAGAGTTCTTTTTACATTCTAGATAAAGGTCCCTTATCAGATATGC
                                                                                                                                                                                                                                                                                          736 TT-TTCAAATATAATCTATAGACTGTCTTTTCAC---TTTCTTGGTGTCCTTTAAAAACAC
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                                                                                                         Gaps
                                                   Score 78; DB 4; Length 75216;
Pred. No. 1.6e-08;
1; Mismatches 161; Indels 15;
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55.1%; Pred. No. 1.1e-07;
cive 1; Mismatches 162; Indels 10;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        Sequence 12055, Application US/09949016
Patent No. 6812339
GENERAL INFORMATION:
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                // LOCATION: (1)...(141454)
// OTHER INFORMATION: n = A,T,C or G
US-09-949-016-12055
                                                   1.8%;
                                              Query Match
Best Local Similarity 56.4
Matches 229; Conservative
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Matches 212; Conservative
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US-09-949-016-14061
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LENGTH: 141454
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Sequence 13379, Application US/09949016

Sequence 13379, Application US/09949016

Patent No. 6812339

TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF

TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF

TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF

CURRENT APPLICATION NUMBER: US/09/949,016

CURRENT FILING DATE: 2000-04-14

PRIOR PLILING DATE: 2000-10-20

PRIOR PLILING DATE: 2000-10-03

PRIOR PLILING DATE: 2000-10-03

PRIOR APPLICATION NUMBER: 60/231,498

PRIOR PLILING DATE: 2000-09-08

NUMBER OF SEQ ID NOS: 207012

SOFTWARE PEBLICS OF Windows Version 4.0

SEQ ID NO 13379
                                                                                                                          14215 ACTANAGGIGTTGATAGICTTTTCATGIGCTAATTTIGCCACCTACACTTCTTCTGATG 144156
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4177 TATTTTTATATTTGTGATCTTACCTTGAATTGAT--ATGTATGTTGTGAATTATGGATCAG 4234
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       3839 ACAAATATTGGGTGTTGTCCAGTATTTTTCCCTTTTTTAACCMTTCCCAATTCGGGTGTGT 3898
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APPLICANT: Goldman, Barry S.
APPLICANT: Hinkle, Gregory J.
APPLICANT: Slater, Steven C.
APPLICANT: Wiegand, Roger C.
TITLE OF INVENTION: Myxococcus xanthus Genome Sequences and Uses Thereof
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           144095 AAGAATTCTTTATACATTCTGGATACCAGTCCTTCAGCAGACATATGATTTAAAATTTTA
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                                                                                     GGTTCTTTTTTCCCCCATACAAGTATCCAGTCATTGTAACACTGTTTATTGAAAGAATT
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                                                                                                                                                                                                                  126723 Trracrircricarrargrircric 126747
                                                                                                                                                                          4295 ATCCTTTCCTCATTAAATTACCTTG 4319
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Matches 172; Conserv
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US-09-902-540-1357/c
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US-09-949-016-13379
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RESULT 13
US-09-949-016-15784/c

i Gequence 15784, Application US/09949016

patent No. 6812339

GENERAL INFORMATION:

i TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED

TITLE OF INVENTION: WITH HUMAN DISBASE, METHODS OF DETECTION AND USES THEREOF

i TITLE OF INVENTION: WITH HUMAN DISBASE, METHODS OF DETECTION AND USES THEREOF

i CURRENT APPLICATION NUMBER: US/09/949,016

CURRENT FILING DATE: 2000-04-14

PRIOR FILING DATE: 2000-10-20

PRIOR PLING DATE: 2000-10-20

PRIOR PLING DATE: 2000-10-03

PRIOR PLING DATE: 2000-10-03

PRIOR FILING DATE: 2000-09-08

NUMBER OF SEQ ID NOS: 207012

SOFTWARE FRALESCO for Windows Version 4.0

SEQ ID NO 15784
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           3959 AGAAATICITIATACATICIAGAIGCAAGICICITGYCGGAIAIACGIAIIGAGAIAITA 4018
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  3899 AGGIGGAIGITICCAITITGGGITTITAATTIGIATAICCCIGAIAGCIATAATIGGGICAT 3958
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        4079 TTGACAAGGTCAAATTTATTTTTTTTTTTTTTGATATTTTTTTCTCTCCAATTTAACCC 4138
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               4139 CAAGATTTCAGATATTCTGCTCTATTATAAACTTTATATTTTTTATATTTGTGATCTAC 4198
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      445
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             3839 ACAAATATTGGGTGTTGTCCAGIATTTTTCCCTTTTTTAACCMTTCCCAATTCGGGTGTGT
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    3779 TTTTTATATTTGTCAATTTATGAATATAATGAATGAGGAGTTCTGGTACCTCCTGTCTTT
                                                                                                                                                                                                                                                                                                                                                                                                                             Length 612;
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                                                                                                                                                                                                                                                                                                                                                                                                                             Score 69.6; DB 4;
Pred. No. 1.4e-07;
1; Mismatches 242;
                                                                                                                                                                                                                                                                                           NAME/KEY: unsure

| LOCATION: (1)..(612)

| OTHER TOPRMATION: unsure at all n locations

US-09-902-540-1357
FILE REFERENCE: 38-10(15849)B
CURRENT APPLICATION NUMBER: US/09/902,540
CURRENT FILING DATE: 2001-07-10
PRIOR APPLICATION NUMBER: 60/217,883
PRIOR FILING DATE: 2000-07-10
NUMBER OF SEQ ID NOS: 16825
SEQ ID NO 1357
                                                                                                                                                                                                                                                                                                                                                                                                                                1.6%;
                                                                                                                                                                                                                   TYPE: DNA ORGANISM: Myxococcus xanthus
                                                                                                                                                                                                                                                                                                                                                                                                                                                      Best Local Similarity 47.9
Matches 224; Conservative
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## Sequence 3149, Application US/09949016
## TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
## TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
## STIER REPERENCE: CLOO1307
## CURRENT APPLICATION NUMBER: 60/241,755
## PRIOR PLIING DATE: 2000-10-20
## PRIOR PLIING DATE: 2000-10-03
## PRIOR PLIING DATE: 2000-10-03
## PRIOR PLIING DATE: 2000-10-03
## PRIOR PLIING DATE: 2000-09-08
## PRIOR FILING DATE: 2000-09-08
## NUMBER OF SEQ ID NOS: 207012
## SOFTWARE: PRAELECE for Windows Version 4.0
64118 TTGTTGTTGTTGAGTTTTAGACATTCCCTATATATTCTGGATATCAATCCCTTATCAGAT 64059
                                                                                                                                                                                                                                                                        53938 ATCCAAGAAATTACTGGCAAATCCAATGTCAAGTTTTGCCCCATGTTTTCTTCAAAG 63879
                                                                                                                                                                                                                                                                                                                                                                 33878 AGTTTTATAGTTGTAGGTCATGGAACCACTCTGAGTTAATTTTTATACACTGTGTTAGGT 63819
                                                                                                                                                                                                                                                                                                                   4116 AT-----TITITICICICCAAITITAACCCCAAGAITICAGATAITCIGCICITAITATATA 4169
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                                                                                        54058 ACAATTTGCAAATATTTTCTCCCATTCTGTGGCTTGCCTTTTTCCTCTACTGATAGTCTT
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56.1%; Pred. No. 4.1e-07;
ive 2; Mismatches 119; Indel8
                                             ATACGTATTGAGATATTACACCTAGTCTGTGGCTTGACTGTTT
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        Query Match
Best Local Similarity 56.11
Matches 169, Conservative
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 RESULT 15
US-09-949-016-31749/c
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US-09-949-016-31749
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LENGTH: 601
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Sequence 12449, Application US/09949016

Patent No. 681239

GENERAL INFORMATION:

APPLICANT: VENTER, J. Craig et al.

TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF

TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF

PILE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF

CURRENT APPLICATION NUMBER: US/09/949,016

PRIOR PELING DATE: 2000-04-14

PRIOR PLING DATE: 2000-10-20

PRIOR PLING DATE: 2000-10-03

PRIOR PLING DATE: 2000-10-03

PRIOR PLING DATE: 2000-10-03

NUMBER OF SEQ ID NOS: 207012

SEQ ID NO 12449

LENGTARE: FastSEQ for Windows Version 4.0

SEQ ID NO 12449

LENGTH: 146039
                                                                                                                                                                                                                                                                                                                                         TIGATGCACACATTITAAGAATTITGGTAAAGICCAAITIGICTATTITTCCTTIGCCAT 35609
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                                                                                                                                                                                                   Length 69709;
                                                                                                                                                                                                   Score 68.2; DB 4; Length 6 Pred. No. 4.9e-06; 1; Mismatches 119; Indels
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1.5%; Score 68.2; DB 4;
Best Local Similarity 56.5%; Pred. No. 7.5e-06;
Matches 170; Conservative 1; Mismatches 119;
                                                           | FEATURE:
| NAME/KEY: misc_feature
| LOCATION: (1)...(69709)
| OTHER INFORMATION: n = A,T,C or G
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OTHER INFORMATION: n = A,T,C or
                                                                                                                                                                                                   Query Match
Best Local Similarity 56.5%;
Matches 170; Conservative
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LOCATION: (1)...(14603)
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US-09-949-016-12449/c
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                   TYPE: DNA
ORGANISM: Human
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LENGTH: 69709
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Search completed: August 30, 2005, 15:39:18 Job time : 684 secs

Sequence Sequence Sequence

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US-09-783-590-3449-

US-09-783-590-3449-

US-09-923-876-5167-

US-09-923-876-5167-

US-09-917-800A-803-

US-09-917-800A-803-

US-09-917-800A-803-

US-09-864-761-11287-

US-09-864-761-11287-

US-09-864-761-11287-

US-09-864-761-11287-

US-09-864-867-425-

US-10-311-455-331-

US-10-85-864-867-425-

US-10-984-867-425-

US-10-786-970A-425-

US-10-786-970A-425-
US-10-146-502-1737
US-10-102-524-532
US-10-242-535A-26197
US-10-085-783A-26197
US-10-085-783A-7179
US-10-085-783A-7179
US-09-918-995-13334
US-09-969-034-1974
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   Sequence 34, Application US/10474495; Publication No. US20040171003A1; GENERAL INFORMATION:
APPLICANT: YOSHIKAWA, YOSHIE et al.; TITLE OF INVENTION: CANCER-ASSOCIATED GENES; FILE REFERENCE: 1422-06.06 PTICKENT FILING DATE: 2003-10-09; PRIOR APPLICATION NUMBER: US/10/474,495; CURRENT FILING DATE: 2001-04-10; PRIOR APPLICATION NUMBER: UP 2001-112039; PRIOR APPLICATION NUMBER: UP 2001-290193; PRIOR PILING DATE: 2001-09-21; NUMBER OF SEQ ID NOS: 264; SOFTWARE: Patentin Ver. 2.1; SEQ ID NO 34
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     ALIGNMENTS
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          NAME/KEY: variation
LOCATION: 553, 597
OTHER INFORMATION: n is a or c or g or t.
                                                                                                                                                                                                                                                                                                                                         6237 15 U

3285 11 U

3285 21 U

3285 21 U

1980090 21

539 14 U

9539 15 U

9539 15 U

10606 15 U

6056 18 U

15548 15 U
     114
6237
6237
3285
     TYPE: DNA
ORGANISM: Homo sapiens
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           US-10-474-495-34
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69.6
69.6
                   507
478.4
478.4
438.2
405.6
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258.4
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239.4
239.4
212.6
191
1167.6
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119.4
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92.4
81.6
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74.6
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72.2
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Sequence 343, App
Sequence 343, App
Sequence 143, App
Sequence 1463, Ap
Sequence 1737, Ap
                                                                                                                            August 30, 2005, 05:17:43 ; Search time 2560 Seconds (without alignments) 11253.603 Million cell updates/sec
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| Cgn2_6/ptodata/2/pubpna/USO6_PUBCOMB.seq:*
| Cgn2_6/ptodata/2/pubpna/USO6_PUBCOMB.seq:*
| Cgn2_6/ptodata/2/pubpna/USO6_PUBCOMB.seq:*
| Cgn2_6/ptodata/2/pubpna/USO6_PUBCOMB.seq:*
| Cgn2_6/ptodata/2/pubpna/USO9_PUBCOMB.seq:*
| Cgn2_6/ptodata/2/pu
                     GenCore version 5.1.6
Copyright (c) 1993 - 2005 Compugen Ltd.
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US-09-22-217-343
US-09-833-263-343
US-10-025-380-343
I US-09-96-034-1463
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US-09-96-935-1737
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                                                                                                                                                                                                                                                                                                                                             7331713 segs, 3271544945 residues
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Maximum Match 100%
Listing first 45 summaries
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Gapop 10.0 , Gapext 1.0
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seg length: 200000000
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Post-processing:

Database

Minimum DB Maximum DB

Title: Perfect score:

Sequence:

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Run on:

Scoring table:

Searched:

Sequence 331, App Sequence 332, App Sequence 425, App Sequence 425, App Sequence 1, Appli Sequence 6815, Ap

Sequence 7179, Ap Sequence 7179, Ap Sequence 13834, A Sequence 1974, Ap Sequence 116439, Sequence 5167, Ap Sequence 8029, Ap Sequence 8727, Ap Sequence 8727, Ap Sequence 8727, Ap Sequence 11287, Ap Seque

Sequence 1369, Ap Sequence 409, App Sequence 52, Appl Sequence 55020, A Sequence 120013, Sequence 12013, Sequence 3, Appl Sequence 73, App Sequence 65, Appl Sequence 673, Appl

Sequence 1, Appli Sequence 2128,

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Length 622; Indela

DB 19;

Score 564.6; DB 19; Pred. No. 1.1e-113; 0; Mismatches 11;

Query Match
Best Local Similarity 97.6%;
Matches 604; Conservative

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                   TTTTGGCTCCGGGAACGTGTGTTGATACAGGAATAACAAAACTCACAGACCTTAATTC
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GENERAL INFORMATION:
APPLICANT: Xu, Jiangchun
APPLICANT: Lodes, Michael J.
APPLICANT: Secriet, Heather
APPLICANT: Benson, Darin R.
APPLICANT: Benson, Darin R.
APPLICANT: Meagher, Madeleine Joy
APPLICANT: Stolk, John A.
APPLICANT: Ming, Yuqiu
APPLICANT: Sinit, Carole Lynn
APPLICANT: Sinit, Carole Lynn
APPLICANT: Sinit, Carole Lynn
APPLICANT: Ming, Ajun
APPLICANT: Ming, Ajun
APPLICANT: Clapper, Jonathan D.
TITLE OF INVENTION: Compounds FOR IMMUNOTHERAPY AND DIAGNOSIS
TITLE OF INVENTION: Compounds FOR IMMUNOTHERAPY
TITLE OF INVENTION: Compounds FOR THEIR USE
FILE REFERENCE: 210121-471C13
CURRENT APPLICATION NUMBER: US/09/922,217
CURRENT APPLICATION NUMBER: US/09/922,217
SUFFMENT FILING DATE: 2001-08-03
NUMBER OF SEQ ID NOS: 1124
SOFTWARE: FESTESEQ for Windows Version 4.0
SEQ ID NO 343
LENGTH: 646
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           Sequence 343, Application US/09922217
Patent No. US20020076414A1
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US-09-922-217-343/c
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Fatent No. US20020110547A1

GENERAL INFORMATION:

APPLICANT: Wang, Ajun

APPLICANT: Stolk, John A.

APPLICANT: Glapper, Jonathan D.

APPLICANT: Manghar, Madeleine J.

APPLICANT: Manghar, Madeleine J.

APPLICANT: Manghar, Madeleine J.

TITLE OF INVENTION: COMPOUNDS FOR IMMUNOTHERAPY AND

TITLE OF INVENTION: DIAGNOSIS OF COLON CANCER AND METHODS FOR THEIR USE

FILE REFERENCE: 2010-1.471C12

CURRENT PLING DATE: 2001-04-10

NUMBER OF SEQ ID NOS: 1093

SEQ ID NOS: 1093

SEQ ID NO 343

LENGTH: 646
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                                        10;
  Length 646;
                                        Indels
DB 9;
Query Match 12.5%; Score 552.2; DB 9; Best Local Similarity 93.0%; Pred. No. 5.8e-111; Matches 609; Conservative 4; Mismatches 32;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          ; TYPE: DNA
; ORGANISM: Homo sapien
US-09-833-263-343
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APPLICANT: Batle, Jon H.
APPLICANT: Carroll, Eddie III
APPLICANT: Carroll, Eddie III
APPLICANT: Dwivedi, Poornima
APPLICANT: Molino, Gary A.
APPLICANT: Thiagalingam, Arunthathi
APPLICANT: Lhiagalingam, Arunthathi
APPLICANT: Lewis, Marcia E.
TITLE OF INVENTION: Nucleic Acid Sequences Differentially
                                                                                                                                                                         DB 13;
                                                                                                                                                                     Query Match
12.5%; Score 552.2; DB 13;
Best Local Similarity 93.0%; Pred. No. 5.8e-111;
Matches 609; Conservative 4; Mismatches 32;
 CURRENT APPLICATION NUMBER: US/10/025,380 URRENT FILING DATE: 2001-12-19 NUMBER OF SEQ ID NOS: 1129 SOFTWARE: PagtSEQ for Windows Version 4.0
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Sequence 1463, Application US/09969034 Publication No. US20040110668A1 GENERAL INFORMATION:
                                                                                                    TYPE: DNA
CORGANISM: Homo sapiens
US-10-025-380-343
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US-09-969-034-1463/c
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LENGTH: 646
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                                                                   3419 AAAAGCTTCTCAGCCTTCCTAGGGAACAGAAATTGGGTGAGCCAATCTGCAATTTCTACT
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 Length 646;
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     DB 9;
Score 552.2; DB 9,
Pred. No. 5.8e-111,
4; Mismatches 32;
12.5%;
al Similarity 93.0%;
609; Conservative
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                                                                                                                                                                                                                                                           12.4%; Score 544.8; DB 11; Length 561; ilarity 99.3%; Pred. No. 2.3e-109; Conservative 0; Mismatches 3; Indels 1;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     Sequence 1737, Application US/09878178
Sequence 1737, Application US/09878178
Patent No. US20020177552A1
GENERAL INFORMATION:
APPLICANT: Jiang, Yuqiu
APPLICANT: Harlocker, Susan L.
APPLICANT: Secrist, Heather
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE THERAPY
TITLE OF INVENTION: AND DIAGNOSIS OF COLON CANCER
TITLE OF INVENTION: Expressed in Cancer Tissue FILE REFRENCE: 1657/1032
CURRENT APPLICATION NUMBER: US/09/969,034
CURRENT FILING DATE: 2001-10-02
PRIOR APPLICATION NUMBER: 60/237,271
PRIOR FILING DATE: 2000-02-10
NUMBER OF SEQ ID NOS: 4494
SOFTWARE: FastSEQ for Windows Version 4.0
SEQ ID NO 1463
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LOCATION: 546
OTHER INFORMATION: n = A,T,C or
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FEATURE:
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Best Local Similarity
Matches 557; Conserv
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US-09-878-178-1737
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US-10-046-935-1737
US-10-046-935-1737

Squence 1737, Application US/10046935

Publication No. US20020156011A1

GRENRAL INFORMATION:
APPLICANT: Jiang, Yuqiu

APPLICANT: Harlocker, Susan L.
APPLICANT: Wang, Aijun

APPLICANT: Scolik, John A.
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE THERAPY

TITLE OF INVENTION: COMPOSITIONS OF COLON CANCER

FILE REFERENCE: 210121:527C1

CURRENT PAPLICATION NUMBER: US/10/046,935

CURRENT FILING DATE: 2002-01-15

NUMBER OF SEQ ID NOS: 2239

SOFTWARE: FastSEQ for Mindows Version 4.0
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Pred. No. 9.5e-108;
1; Mismatches 2;
                       US/09/878,178
FILE REFERENCE: 210121.527
CURRENT APPLICATION WUMBER: US/09/878,
CURRENT FILING DATE: 2001-06-08
NUMBER OF SEQ ID NOS: 2237
SOFTWARE: FastSEQ for Windows Version
SEQ ID NO 1337
LENGTH: 541
                                                                                                                                                                                                                                                   12.2%;
                                                                                                                                                                                                                                                                                                  538; Conservative
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; ORGANISM: Homo sapien
US-09-878-178-1737
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Best Local Similarity
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3610 TGAAATGGAGTCATTTGAGTCTYTTAATAGCCATGTATCATAATTACCAAGTGAAGCTGG 3669
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Publication No. US20030109434A1
GENERAL INFORMATION:
APPLICANT: Algate, Paul A.
APPLICANT: Mannion, Jane
APPLICANT: Garden, Brian
APPLICANT: Harlocker, Susan L.
TITLE OF INVENTION: COMPOSITIONS AND DIAGNOSIS OF KIDNEY CANCER
TITLE OF INVENTION: THERAPY AND DIAGNOSIS OF KIDNEY CANCER
FILE REFERENCE: 210121.572
CURRENT APPLICATION NUMBER: US/10/102,524
CURRENT FILING DATE: 2002-03-19
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             Score 537.4; DB 14;
Pred. No. 9.5e-108;
1; Mismatches 2;
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Best Local Similarity 93.6%; Pred. No. 4.9e-101;
Matches 559; Conservative 4; Mismatches 24;
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SOFTWARE: FastSEQ for Windows Version 4.0
SEQ ID NO 532
                Query Match
Best Local Similarity 99.4%;
Matches 538; Conservative
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TYPE: DNA
ORGANISM: Homo sapiens
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APPLICANT: Harlocker, Susan L.
APPLICANT: Harlocker, Susan L.
APPLICANT: Secrist, Heather
APPLICANT: Stolk, John A.
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE THERAPY
TITLE OF INVENTION: COMPOSITIONS OF COLON CANCER
FILE REFERENCE: 210121.52762
CURRENT APPLICATION NUMBER: US/10/146,502
CURRENT APPLICATION NUMBER: 2231
NUMBER OF SEQ ID MOS: 2241
SOFTWARE: FastSEQ for Windows Version 4.0
                                                                                                         DB 13; Length 541;
                                                                                                                                            2; Indels
                                                                                                       12.2%; Score 537.4; DB 1399.4%; Pred. No. 9.5e-108
                                                                                                                                        1; Mismatches
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Publication No. US20030069180A1
GENERAL INFORMATION:
                                                                                                                      Best Local Similarity 99.4 Matches 538; Conservative
                                 TYPE: DNA
CORGANISM: Homo sapiens
US-10-046-935-1737
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; ORGANISM: Homo sapiens
US-10-146-502-1737
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US-10-146-502-1737
SEQ ID NO 1737
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LENGTH: 541
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Best Local Similarity 99.6%; Pred. No. 8.7e-95; Matches 479; Conservative 0; Mismatches 2;
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US-10-24-535A-26197

Sequence 26197, Application US/10242535A

Sequence 26197, Application Wo/US20040013663A1

GENERAL INFORMATION:

APPLICANT: ChondroGene Inc.

TITLE OF INVENTION: Compositions and Methods Relating to Osteoarthritis

TITLE OF INVENTION: Compositions and Methods Relating to Osteoarthritis

TITLE OF INVENTION: Compositions and Methods Relating to Osteoarthritis

CURRENT APPLICATION NUMBER: US/10/242,535A

CURRENT FILING DATE: 2002-09-12

PRIOR PPLICATION NUMBER: US 60/305,340

PRIOR FILING DATE: 2001-07-13

PRIOR PILING DATE: 2001-03-12

PRIOR SPOILOATION NUMBER: US 60/275,017

PRIOR SPOILOATION NUMBER: US 60/271,955

NUMBER OF SEQ ID NOS: 58994

SOFTWARE: PatentIn version 3.2
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 3419 AAAAGCTTCTCAGCCTTCCTAGGGAACAGAAATTGGGTGAGCCAATCTGCAATTTCTACT 3478
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                                    1 AAAAGCTTCTCAGCCTTCCTAGGGAACAGAAATTGGGTGAGCCAATCTGCAATTTCTACT
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LOCATION: (31)...(31)
OTHER INFORMATION: n is a, c, g,
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LENGTH: 481
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10.9%; Score 478.4; DB 17; Length 481;

Query Match

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RESULT 11
US-10-085-783A-26197
; Sequence 26197, Application US/10085783A
; Sequence 26197, Application No. US20040037841A1
; Publication No. US20040037841A1
; GENERAL INFORMATION:
; APPLICANT: ChondroGene Inc.
; APPLICANT: ChondroGene Inc.
; APPLICANT: Liew, C.C.
; TITLE OF INVENTION: Compositions and Methods Relatiing to Osteoarthritis
; FILE REPERROR: 4231/2002
; CURRENT FILING DATE: 2002-02-28
; CURRENT FILING DATE: 2001-07-13
; PRIOR FILING DATE: 2001-07-13
; PRIOR FILING DATE: 2001-07-13
; PRIOR FILING DATE: 2001-03-12
; PRIOR FILING DATE: 2001-03-12
; PRIOR FILING DATE: 2001-03-12
; PRIOR FILING DATE: 2001-03-13
; PRIOR FILING DATE: 2001-03-13
; SETON FILING DATE: 2001-03-14
; SEQ ID NOS: 58994
; SEQ ID NO 26197
; LENGTH: 481
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                                                3126 ATGTACCTGTGATCCAGCAAGAAGGGAGTTCCAGTCAAGAGTCACTACAACTGATTAGTT
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Pred. No. 8.7e-95;
0; Mismatches 2; Indels 0;
Indels
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Best Local Similarity 99.6%;
Matches 479; Conservative
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RESULT 13
US-10-085-783A-7179
is Sequence 7179, Application US/10085783A
j Publication No. US20040037841A1
general information:
j APPLICANT: Chandrodene Inc.
j APPLICANT: Chandrodene Inc.
j TITLE OF INVENTION: Compositions and Methods Relating to Osteoarthritis
fILE REFERENCE: 431/2002
j CURRENT APPLICATION NUMBER: US/10/085,783A
j CURRENT PILING DATE: 2002-02-28
j PRIOR FILING DATE: 2001-07-13
j PRIOR FILING DATE: 2001-07-13
j PRIOR FILING DATE: 2001-03-12
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                                                                    TCTGCAATTTCTACTACAGGCATTGAGACCAGTTAGATTTTGAAATTATTAGAGAGTT
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Pred. No. 5.9e-86;
2; Mismatches 0; Indels
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Best Local Similarity 99.5%;
Matches 437; Conservative
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US-10-085-783A-7179
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LENGTH: 439
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; Sequence 7179, Application US/10242535A
; Publication No. US20040013663A1
; GENERAL INFORMATION:
    APPLICANT: ChondroGene Inc.
; APPLICANT: ChondroGene Inc.
; APPLICANT: ChondroGene Inc.
; TITLE COF INVENTION: Compositions and Methods Relatiing to Osteoarthritis
FILE REFERENCE: 4231/2005
; CURRENT APPLICATION NUMBER: US/10/242,535A
; CURRENT APPLICATION NUMBER: US 10/085,783
; PRIOR APPLICATION NUMBER: US 60/305,340
; PRIOR PILING DATE: 2001-03-12
; PRIOR PILING DATE: 2001-03-12
; PRIOR APPLICATION NUMBER: US 60/275,017
; PRIOR APPLICATION NUMBER: US 60/271,955
; NUMBER OF SEQ ID NOS: 58994
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 7179
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                                                                                                                                 61 GITIAGAGAATGAGAATGGAACAGTGAGGAATGGAGGCCATATITCCATGACTTCCTT
                                                                                                                                                                                                                                                                 TTGCAGGTGGAACTCCAGCTGCAAGGGAGTTAGGGAAATGAAGGTCTTTTTAAAAGCT
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  3126 ATGTACCTGTGATCCAGCAAGAAGGGAGTTCCAGTCAAGAGTCACTACAACTGATTAGTT
                         1 ATGTACCTGTGATCCAGCAAGTAGGGAGTTNCAGTCAAGAGTCACTACAACTGATTAGTT
                                                                                        GTTTAGAGAATGAAATGGAACAGTGAGGAATGGAGGCCATATTTCCATGACTTCCCTT
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Best Local Similarity 99.5%; Pred. No. 5.9e-86;
Matches 437; Conservative 2; Mismatches 0
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US-10-242-535A-7179
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US-10-242-535A-7179
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         413 INGGITAANAINAAGAICCCAAACIAICAGCITICNCCAAATTAAANIGGCCAAGNIAIA
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 1433 CCGTCTACCAAAAAAGGGAAAGGGATGAAGAAATCAAGAAATGTCTTCATGCAAATCA
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            173 IGGAAAATAAGGAGCAGCAINTAINIGAGAATGAGCCTGNGGACNCAAACTAGGNAAAT
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                                                                                                                                                                                                                                   APPLICANT: Agle, Jon H.
APPLICANT: Agle, Jon H.
APPLICANT: Carroll, Eddie III
APPLICANT: Carroll, Eddie III
APPLICANT: Catino, Theodore J.
APPLICANT: Dwivedi, Poornima
APPLICANT: Molino, Gary A.
APPLICANT: Thiagalingam, Arunthathi
APPLICANT: Molino, Gary A.
TITLE OF INVENTION: Expressed in Cancer Tissue
FILE REFERENCE: 1657/1032
CURRENT APPLICATION NUMBER: US/09/969,034
CURRENT FILING DATE: 2001-10-02
PRIOR APPLICATION NUMBER: 60/237,271
PRIOR APPLICATION NUMBER: 60/237,271
SEQ ID NOS: 4494
SOFTWARE: FastSEQ for Windows Version 4.0
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331,
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315,
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Pred. No. 3.8e-69;
0; Mismatches 74;
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303,
417,
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301,
412,
                                                                                                                                     US-09-969-034-1974/c
; Sequence 1974, Application US/09969034
; Publication No. US20040110668A1
; GENERAL INFORMATION:
; APPLICANT: Burgess, Christopher C.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            | NAME/KEY: miso_feature
| LOCATION: 36, 41, 58, 70, 85, 118,
| LOCATION: 203, 211, 217, 247, 289,
| LOCATION: 358, 367, 378, 402, 405,
| LOCATION: 444, 470, 475, 477
| COTHER INFORMATION: n = A,T,C or G
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Best Local Similarity 83.8
Matches 383; Conservative
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      TYPE: DNA
ORGANISM: Homo sapiens
                 CT 3359
                                                       CT 468
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                                                                               3644 GTATCATAATTACCAAGTGAAGCTGGTGGAACATATGGTCTCCATTTTACAGTTAAAGGAA 3703
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                                                                                                                                                                                                                                                                                                                                                                            SQUENCE 13834, Application US/09918995

PUBLICATION NO. US20020073623A1

GENERAL INFORMATION:

APPLICANT: Hyseq. Inc.

TITLE OF INVENTION: NOVEL NUCLEIC ACID SEQUENCES OBTAINED

TITLE OF INVENTION: FROM VARIOUS cDNA LIBRARIES

FILE REFERENCE: 20411-756

CURRENT APPLICATION NUMBER: US/09/918,995

CURRENT FILING DATE: 1299-01-20

NUMBER OF SEQ ID NOS: 38054

SOFTHARE: PSECSEQ for Windows Version 3.0

LENGTH: 468
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TAGAATTGACAGGCCATATTAGTTGATGAAATGGAGTCATT
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                                                                                                                                                                                                                                                3764 GCCCTACTATAGCAGTTTT 3782
                                                                                                                                                                                                                                                                     421 GCCCTACTATAGCAGTTTT 439
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 ; LOCATION: (1)...(468); OTHER INFORMATION: n = A,T,C or US-09-918-995-13834
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US-09-918-995-13834
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GenCore version 5.1.6  Copyright (c) 1993 - 2005 Compugen Ltd.  OM nucleic - nucleic search, using sw model  Run on: August 30, 2005, 05:15:28; Search time 13109 Seconds (without alignments) 12784.879 Million cell updates/sec Title: US-09-837-138-1 Perfect score: 4403 Sequence: 1 ttcggcacgaggcgcggttgaccgcggtggagctccagct 4403 Scoring table: Capop 10.0, Gapext 1.0 Searched: 34239544 seqs, 19032134700 residues Total number of hits satisfying chosen parameters: 68479088	length: 0 length: 20000000	Post-proceesing Minimum Match 0%  Database : EST;  Ligting first 45 summaries  Listing first 45 summaries  EST;  2

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REMARK NIH-MGC Project URL: http://mgc.nci.nih.gov COMMENT Contact: MGC help desk Email: cgapbe-remail.nih.gov Tissue Procurement: Miklos Palkovits, M.D., Ph.D. Tissue Procurement: Miklos Palkovits, M.D., Ph.D. CDNA Library Preparation: Michael J. Brownstein (NHGRI) & Shiraki Toshiyuki and Piero Carninci (RIKEN) CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL) DNA Sequencing by: The I.M.A.G.E. Consortium (LLNL) DNA Sequencing by: Sequencing Group at the Stanford Human Genome Center, Stanford University School of Medicine, Stanford, CA 94305 Web site: Contact: (Dickson, Mark) mcdepaxil.stanford.edu Dickson, M., Schmutz, J., Grimwood, J., Rodriquez, A., and Myers, R. M.  Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LNL at: http://image.llnl.gov Series: IRAK Plate: 63 Row: a Column: 1 This clone was selected for full length sequencing because it passed the following selection criteria: matched mkNA gi: 6996019 This clone has the following problem: frame shifted.    1.4550		Qy         263 AATTCRAGTACGTACTTTATATAGGAAAAATGCAGATGCGAATTTCCCGAACT         322
0	BC040519 BC040519 BC040519 BC040519 BC040519 LOCUS BC040519 LOCUS DEPINITION Clone IMAGE:4791579), containing frame-shift errors. Clone IMAGE:4791579), containing frame-shift errors. BC040519.1 G1:26996492 KENSION BC040519.1 G1:26996492 HTC BC040519.1 G1:26996492 HTC BC040519.1 G1:26996492 REPRINCE HOWO sapiens HOW Sapiens RaryOta; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; BaryOta; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; BaryOta; Metazoa; Primates; Catarrhini; Hominidae; Homo. REFERENCE Rausberg, R.L., Feingold, E.A., Grouse, L.H., Derge, J.G., Altschul, S.F., Zeeberg, B., Buerow, K.H., Schaefer, C.F., Bhat, M.K., HOPKINS, R.F., Jordan, H., Moore, T., Max, S.I., Wang, J., Hsieh, F., Diatchenko, L., Marusina, K., Farmer, A.A., Rubin, G.M., Hong, L., Scheetz, T.E., Brownstein, M.J., Usdin, T.B., Toshiyuki, S., Carninci, P., Prange, C., Raha, S.S., Loquellano, N.A., Peters, G.J., McKernan, K.J., Malek, J.A., Guarathe, P.H., Richards, S., Worley, K.C., Hale, S., Garcia, A.M., Gay, L.J., Hulkk, S.W., Villalon, D.K., Mullay, S., Ketteman, M., Madan, A., Rodrigues, S., Fahev, J., Helton, E., Ketteman, M., Madan, A., Rodrigues, S.,	Sanchez, A., Whiting, M., Madan, A., Young, A.C., Shevchenko, Y., Bouffard, G.C., Blakesley, R.W., Touchman, J.W., Green, E.D., Dickson, M.C., Redriguez, A.C., Touchman, J.W., Green, E.D., Butterfield, Y.S., Krzywinski, M.I., Schaleka, U., Smailus, D.E., Schnerch, A., Schein, J.E., Jones, S.J. and Marra, M.A.  TITLE Generation and initial analysis of more than 15,000 full-length human and mouse cDNA sequences Cournal Proc. Natl. Acad. Sci. U.S.A. 99 (26), 16899-16903 (2002) REDERENCE 2 (bases 1 to 4550) AUTHORS Strausberg, R. TITLE Strausberg, R. TITLE Gene Collection (MGC), Cancer Genomics Office, National Cancer Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590, USA

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Mamalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 2265)
Clark,A.G., Glanowski,S., Nielson,R., Thomas,P., Kejariwal,A.,
Todd,M.A., Tanenbaum,D.M., Civello,D.R., Lu,F., Murphy,B.,
Pertera,S., Wang,G., Zheng,X.H., White,T.J., Sninsky,J.J.,
Adams,M.D. and Cargill,M.
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                                        Clone distribution: MGC clone distribution information can be four through the I.M.A.G.E. Consortium/LLNL at: http://image.llnl.gov Series: IRAL Plate: 16 Row: j Column: 6 This clone was selected for full length sequencing because it passed the following selection criteria: matched mRNA gi: 3098674 This clone has the following problem: frame shifted.
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Matches 1476;
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933 TGGATATGCTCCAAAGGCAAGGTCTTAGACCTATTCCTGAAGCAGAAATTGGATTGGCGG 992
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                                                                                                                                         Length 1460;
                                                                                                                                                                           Indels
 /db_xref="taxon:9606"
/clone="MAGE:4104186"
/clone="type="Brain, glioblastoma"
/clone lib="wlH MGC_57"
/lab_host="DH108"
/note="Vector: pDNR-LIB"
                                                                                                                                                                           10;
                                                                                                                                         Score 1427; DB 3;
Pred. No. 6.4e-295;
                                                                                                                                                                          0; Mismatches
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99.3%;
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                                                                                                                                                         Similarity
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Best Local Simil
Matches 1433; C
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Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

    (bases 1 to 1460)

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This clone was selected for full length sequencing because it passed the following selection criteria: matched mRNA gi: 6996019
This clone has the following problem: frame shifted.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Direct Submission
Submitted (05-NOV-2001) National Institutes of Health, Mammalian
Gene Collection (MGC), Cancer Genomics Office, National Cancer
Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    Contact: MGC help desk
Email: cgapbs-rémail.nih.gov
Tissue Procurement: ATC
CDNA Library Preparation: CLONTECH Laboratories, Inc.
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Institute for Systems Biology
http://www.systemsbiology.org
contact: amadan@systemsbiology.org
Anup Madan, Jessica Fahey, Brin Helton, Mark Ketteman, Anuradha
Madan, Stephanie Rodrigues, Amy Sanchez and Michelle Whiting
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   1460 bp mRNA linear HTC 09-NOV-:
Homo sapiens, Similar to Nijmegen breakage syndrome 1 (nibrin),
Clone IMAGE:4104186, mRNA.
GGGAAGAAACGTGAACTCAAGGAAGACTCACTATGGTCAGCTAAAGAAATATCTAACAAT
                                                                         GGGAAGAACGTGAACTCAAGGACTCACTATGGTCAGCTAAAGAAATATCTNNNNN
                                                                                                                    GACAAACTTCAGGATGATAGTGAGATGCTTCCAAAAAAGCTGTTATTGACTGAATTTAGA
                                                                                                                                                        TCACTGGTGATTAAAAACTCTACTTCCAGAAATCCGTCTGGCATAAATGATTATGGT
                                                                                                                                                                                                                          CAACTAAAAATTTCAAGAATTCAAAAAGGTCACATATCCTGGAGCAGGAAAACTTCCA
                                                                                                                                                                                                                                                                                                                              CACATCATTGGAGGATCAGATCTAATAGCTCATGCTCGAAAGAATACAGAACTAGAA
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      /organism="Homo sapiens"
/mol_type="mRNA"
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Homo sapiens
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TITLE
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/db_xref="taxon:10090"
/dclone="lthAGE:3602503"
/tissue type="wammary tumor. C3(1)-Tag model. Infiltrating ductal carcinoma. 5 month old virgin mouse."
/clone lib="NCI CAAP_Mam6"
/lab host="DHIOI CAAP_MAF6"
/note="Vector: pCMV-SPORT6"
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LIML at: http://image.llnl.gov Series: IRAK Plate: 17 Row: g Column: 3 This clone was selected for full length sequencing because it passed the following selection criteria: Similarity but not
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       276
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                                                                                                                                                           Submitted (27-AUG-2001) National Institutes of Health, Mammalian Gene Collection (MGC), Cancer Genomics Office, National Cancer Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590,
                                                                                                                                                                                                                                                                                                                                                                                                                                                         Sequencing Center Center Center code: BCM-HGSC Web site: http://www.hgsc.bcm.tmc.edu/cdna/
Contact: amg@bcm.tmc.edu
Gunaratne, P.H., Garcia, A.M., Lu, X., Hulyk, S.W., Loulseged, Kowis, C.R., Sneed, A.J., Martin, R.G., Muzny, D.M., Nanavati, A.N., Gibbs, R.A.
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                                                                                                                                                                                                                                                                       NIH-MGC Project URL: http://mgc.nci.nih.gov
Contact: MGC help desk
Email: cgapbs-r@mail.nih.gov
Tissue Procurement: Jeffrey Green M.D.
cDNA Library Preparation: Life Technologies, Inc.
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Baylor College of Medicine Human Genome
  Natl. Acad. Sci. U.S.A. 99 (26), 16899-16903 (2002)
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      identity to protein.
This clone has the following problem: retained intron.
Location/Qualifiers
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/strain="FVB/N"
                                                                                    to 1792)
                                                                                                                                          Direct Submission
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Strausberg, R.
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Best Local Similarity
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JOURNAL
MEDLINE
PUBMED
REFERENCE
AUTHORS
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COMMENT
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Bukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Bukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Bukaryota; Metazoa; Chordata; Sciurognathi; Muridae; Murinae; Mus.

I (bases 1 to 1792)

Strausberg, R.L., Feingold, E.A., Grouse, L.H., Derge, J.G.,

Altschul, S.F., Zeeberg, B., Buetow, K.H., Schemmen, C.M., Schuler, G.D.,

Altschul, S.F., Jordan, H., Morce, T., Mang, J., Hsieh, F.,

Diatchenko, L., Marusina, K., Farmer, A.A., Rubin, G.M., Hong, L.,

Stapleron, M., Soarses, M., B., Bonaldo, M.F., Casavant, T.L.,

Scheetz, T.E., Brownstein, M.J., Usdin, T.B., Toshiyuki, S.,

Carninci, P., Prange, C., Raha, S.S., Loquellano, N.A., Peters, G.J.,

Abramson, R.D., Mullahy, S.J., Bosak, S.A., McEwan, P.J.,

McKernan, K.J., Malek, J.A., Gunatane, P. H., Richards, S.,

Worley, K.C., Hale, S., Garcia, A.M., Gay, L.J., Hulyk, S.W.,

Villalon, D.K., Muzny, D.M., Sodergren, E.J., Lu, X., Gibbs, R.A.,

Bouffard, G.G., Blakesley, R.W., Touchman, J.W., Green, E.D.,

Butterfield, Y.S., Krzywinski, M.I., Shalska, U., Scheuchenko, Y.,

Butterfield, Y.S., Krzywinski, M.I., Shalska, U., Sanailus, D.E.,

Generation and initial analysis of more than 15,000 full-length
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                                                                                                                         965 TGATTTTCATGACTACAAAGAATTACTGTGATCCTCAGGGCCATCCCAGTACAGGTTAA 1024
                                                                                                                                                                                             AGACAACAACTCCAGGACCAAAGCCTTTCACAAGGGGTGTCAGTTGAAAAACTAATGC 1112
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GAATGCTTTCACAAGACGCACCCACTGTAAAGGAGTCCTGCAAAACAAGCTCTAATAATA
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        CAGTTTGAGTCAAACAGATGAAATTCCTACATTAACAATAAAAGATAATTCTAAGTATGG
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                                                                       318 AACCTTTGTTAATGAAGAAAAAATGCAGACTGGTCTTTCCTGCACGTTGAAGACAGGAGA
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                                              TACCTTTGTTAATGAGGAAAAATGCAGAATGGCTTTTCCCGAACTTTGAAGTCGGGGGA
                                                                                                                                                                                                                                                   438 TTGTTCTTCTTGTTTAGATGTCTCTGGGAAAACTGTTTTAAATCAAGCTATTTTACAGCT
                                                                                                                                                                                                                                                                                           TGGAGGATTTACTGTAAACAATTGGACAGAAGAATGCACTCACCTTGTCATGGTATCAGT
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Katoh, H., Kawai, J., Kojima, Y., Kondo, S., Konno, H., Kouda, M., Koya, S., Kurihara, C., Matsuyama, T., Miyazaki, A., Murata, M., Nakamiza, M., Nakamiza, M., Namazaki, R., Ohno, M., Ohsato, N., Okazaki, Y., Saito, R., Saitoh, H., Sakai, C., Sakai, K., Sakazume, N., Sano, H., Sasaki, D., Shibata, K., Shinagawa, A., Shiraki, T., Sogabe, Y., Tagawai, M., Tagawa, A., Takahashi, F., Takaku-Akahira, S., Takeda, Y., Tanaka, T., Tagawa, A., Toya, T., Yasunishi, A., Direct Submission and Hayashizaki, Y.

Direct Submission Noshihide Hayashizaki, The Institute of Physical and Chemical Research (RIKEN), Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center (GSC), RIKEN Yokohama Institute; 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan (E-mail:genome-res@gsc.riken.jp, URL:http://genome-gsc.riken.jp/, Tel:81-45-503-9222,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         /protein_ide="BAC27610.1"
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QNATLDREADTSSVGGNDIELNRKSPDRKPLPTETLRP"
                                                                                                                                                                                                                                                                                                                          CDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in RIKEN.

Division of Experimental Animal Research in Riken contributed to prepare mouse tissues.

Please visit our web site for further details.

URL:http://fantom.gsc.riken.jp/.

URL:http://fantom.gsc.riken.jp/.

Location/Qualifiers

1. .1731

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GB|NM_013752, evidence: BLASTN,
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Mismatches
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Contact: Genoscope
Contact: Genoscope
Genoscope - Centre National de Sequencage
2 rue Gaston Cremieux, CP 5706 - 91057 EVRY cedex - FRANCE
Email: seqref@genoscope.cns.fr, Web: www.genoscope.cns.fr
1st strand cDNA was primed with a Noil-oligo(dI) primer. Five prime
end enriched, double-strand cDNA was digested with Not I and cloned
into the Not I and EcoRV sites of the pCMVSPORT 6 vector. Library
was not normalized. Library was constructed by Life Technologies, a
division of Invitrogen.
This sequence belongs to sequence cluster 4762.r
For more information about this cluster, see
http://www.genoscope.cns.fr/cdna?e=CSOAM009BH06QPl&c=4762.r.
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/mol_type="mRNA"
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/note="Organ: liver; Vector: pCMVSPORT_6; lst strand cDNA
was primed with a NotI-oligo(dT) primer. Five prime end
enriched, double-strand cDNA was digested with Not I and
cloned into the Not I and ECORV sites of the pCMVSPORT 6
vector. Library was not normalized."
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BX405940 Homo sapiens FETAL LIVER Homo sapiens cDNA clone CSD0009Y012 5-PRIME, mRNA sequence.
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|GACAGGGAAGCCGACACCTCATCTGTGGGTGGGATGGACATAGAACTCAACAGGAAGAG 1691
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Li, W.B., Gruber, C., Jessee, J. and Polayes, D.
Full-length cDNA libraries and normalization
Unpublished (2001)
                                               GAAATTTCCTGTTGCAAATAAAAATAAGGATTGGACTTCTCAGCAGCAGCAGCAG
                                                                                                              CATCAAAAACTACTTCCAGCCATGCACCAGAAAAAGGGAAAGGGATGAAGACAACCCAGA
                                                                                                                                                1474 AATGTCTTCATGCAAATCAGCAAGAATAGAAACGTCTTGTTCTCTTTTAGAACAACACA
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Pred. No. 8.8e-185;
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Eukaryota, Metazoa; Chordata; Craniata; Vertebrata; Buteleostomi; Bukaryota; Metazoa; Chordata; Craniata; Vertebrata; Buteleostomi; Mammalia; Butheria; Primates; Catarrhini; Hominidae; Homo.

I (bases 1 to 1163)

S NIH-MGC http://mgc.nci.nih.gov/.

National Institutes of Health, Mammalian Gene Collection (MGC)

LOnpublished (1999)

LOntact: Robert Strausberg, Ph.D.

Emal: cgapbs-r@mail.nih.gov

Tissue Procurement: ATCC

CDNA Library Preparation: Life Technologies, Inc.

CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LINL)

DNA Sequencing by: Agencourt Bioscience Corporation

Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LINL at: http://image.llnl.gov

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  TMATTACCAAGTGAAGCTGGTGGAACATATGGTCTCCAAMTCACAGTTAAGGAACATAAT 603
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For more information about this cluster, see
http://www.genoscope.cns.fr/cdna?s=CSOAM009BH06NP1&c=4762.r.
Location/Qualifiers
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/note="See 'Creation of Genome-wide Protein Expression
Libraries using Random Activation of Gene Expression,
Nature Biotechnology, in press. Note that even though the
cell type indicated is HT1080, since a random activation
method was used, these sequence tags are not necessarily
expressed in HT1080 under normal circumstances."
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Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
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RST1768 Athersys RAGE Library Homo sapiens CDNA, mRNA sequence.
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3201 Carnegle Ave, Cleveland,
Tel: 216 431 9900
Fax: 216 361 9596
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Location/Qualifiers
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REFERENCE 2 (Dases 1 to 1803)  AUTHORS Clark, A.G., Glanowski, S., Nielson, R., Thomas, P., Kejariwal, A., Tanenbaum, D.M., Civello, D.R., Lu, F., Murphy, B., Ferriera, S., Wang, G., Zheng, X.H., White, T.J., Sninsky, J.J., Adama, M.D. and Cargill, M.  TITLE Adama, M.D. and Cargill, M.  TITLE Submitted (16-NOV-2003) Celera Genomics, 45 West Gude Drive, Borect Submission JOURNAL Submitted (16-NOV-2003) Celera Genomics, 45 West Gude Drive, Rockville, MD 20850, USA  COMMENT This sequence was made by sequencing genomic exons and ordering them based on alignment.  FEATURES  Location/Qualifiers  Location/Qualifiers  Location/Qualifiers  Ab zref="taxon:10090"  /db zref="taxon:10090"  /locus_tag="NBS1"  /locus_tag="HCM5756"	18.7%;   Score 822.2;   DB 9;   Length 1803;	Db 121 ACCTTTGGGGTGTTTGGAAGTTAATTCAGNNNNNNNNNNN	
	Oy         1021         TGATCCTCAGGGCCATCCCAGTACAGGATTAAAGACAACAACTCCAGGACCATTC         1080           Db         501         TGATCCTCAGGGCCATCCCAGTACAGGATTAAAGACAACACCCAGGACCAAGCCTTTC         560           Oy         1081         ACAAGGCGTGTCAGTTGATGAAAACTAATGCCAAGCGCCCCAGTGAACACTACAACATA         1140           Db         561         ACAAGGCGTGTCAGTTGATGAAAACTAATGCCAAGCGCCCCAGTGAACACTACAACATA         620           Oy         1141         CGTAGCTGACACAGAATCAGAGCAGTTACATGGGATTTGAGTGAAAGGCCAAAAGA         1200           Db         621         CGTAGCTGACACAGAATCAGAGCAGATACATGGGATTTGAGTGAAAGGCCCAAAAGA         680           Oy         1201         AATCAAAGTCCCAAAATGGAACAAAAATTCAGAATGCTTTCACAAGAGCCCACTGT         1260           Oy         1201         HIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIII	QY         1261 AAAGGAGTCCTGCAAAACAAGCTCTAATAATAATAATAGTATGGTATCAAATACTTTGGCTAA 1320           Db         741 NAAGGAGTCCTGCANAATAAGCTCTAAT-ATAATAGTATGGGAATACTCTGGCTAA 799           QY         1321 GATGAGAATCCCAAACTATCACCTTCACCAACTAAATTGCCAAGTATAAAAGTAA 1380           Db         800 GATGAGAATNCCAAACTATCACCAACTTAATTGGCCAAGTATAAAAGTA 859           QY         1381 AGATAGGGCTTCTCAGCAGCAGCAGCAACTCCATCAACTACTTCAGCCGTTAC 1440           Db         860 AGATAGGGCTTCTCAGCAGCAGACCAACTCCATCAGAACTACTTCAGGCGTTAC 919           QY         1441 CAAAAAA 1447           Db         920 CACAAAA 926	RESULT 12 AV416036 AV416036 LOCUS LOCUS Mus musculus NBS1 gene, VIRTUAL TRANSCRIPT, partial sequence, genomic survey sequence. ACCESSION AV416036 AVA16036 A

Insert Length: 785 Std Error: 0.00.  Location/Qualifiers  1. 785  1. 785    Arganism="Homo sapiens"     Ab xref="taxon:9606"     Ab xref="taxon:96	Query Match         17.5%;         Score 771.4;         DB 7;         Length 785;           Best Local Similarity         99.7%;         Pred. No. 2.2e-154;         Matches 783;         Conservative         0;         Mismatches         1;         Indels         1;         Gaps         1;           Qy         469         TGTAAACAATTGGACAGAAGAAGAAGCACTCACCTTGTCATGGTATCAGGAAAGTTACCAT         528         528           Dh         1         TGTAAACAATTGGAAGAAGAAGAAGCATCACCTTGTCATGGTATCAGGAAATTACCAT         60	529   TARARCATATATGCCACTCCATTGTGGACGTCCAATTGTAAAGCCACAATTTTACTGA	Oy 649 TCTTGATGAACCATCTATTGGAAGTAAAAATGTTGATCTGTCAGGACGGGGGGGG	Qy 709 ACADATCTTCADAGGADADCATTTATATTTTGADTGCCDACGGGTADGADATTGGG 768	Oy 769 TTCCGCAGTTGTCTTTGGAGGGGGAAGCTAGGTTGATAACAGAAGAGAATGAAGAA 828 	829 ACATAATITCTTITIGGCTCCGGGAACGIGTGTTGTTGATACAGGAATAACAAACTCACA	0.0   0.0
	Oy         1184 AGTGAAAGGCCAAAAGAAATCAAAAGTCTCCAAAATGGAACAAAAATTCAGAATGCTTTCA 1243           Db         961 AGTGAAAGACCAGAAGTAAAGATCCCTGGACTGGAACAAAGCTCTAGGAAACTTTCA 1020           Oy         1244 CAAGACGCACCCACTGTAAAAGATCCTGCAAAACAACTCTAATAATAATAATAGTA 1303           Db         1021 CAAGAAACAATTAAAGAAGACCCCTAAACCTAAACCTAAACTCAAAACTAAACTAACAATTCAAAAACTAACAAC	1304   TCAMATTACTTTGGCTAAGATCCCAAACTATCAGCTTTCACCAACTAAATTGCCA	A-8	Oy 1484 TGCAAATCAGCAAGAATAGAAACGTCTTGTTCTTTTAGAACAAACA	OY 1544 CCCTCATTGTGGAAAAATAAGGGCAGCATCTATCTGAGAATGAGCCTGTGGACACAAAC 1603	Qy 1604 TCAGACA 1610             Db 1375 GCCGACA 1381	CN304420 LOCUS LOC

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1 (Bases 1 to 789)
Harrington, J.J., Sherf, B., Rundlett, S., Jackson, P.D., Perry, R.,
Gain, S., Leventhal, C., Thornton, M., Ramachandran, R.,
Mhittington, J., Lerner, L., Costanzo, D., McBlligott, K., Boozer, S.,
Mys, R., Smith, B., Veloso, N., Klika, A., Hess, J., Cothren, K., Lo, K.,
Offenbacher, J., Danzig, J. and Ducar, M.
Creation of gene expression libraries using random activation of gene expression.
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2373264 Athersys RAGE Library Homo sapiens cDNA, mRNA sequence.
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597 AAAATATCTCAAGAAAATGAAATTGGGAAGAACGTGAACTCAAGGAAGACTCCACTATGG
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21227151
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Athersys, Inc.
3201 Carnegie Ave, Cleveland, Ol
Tel: 216 431 9900
Fax: 216 361 9596
Email: scain@athersys.com
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           Email: paul-mccray@ulowa.edu
Tissue Procurement: Dr. M. J. Welsh, University of Iowa
CDNA Library preparation: Dr. M. Bento Soares, University of Iowa
CDNA Library Arrayed by: Dr. M. Bento Soares, University of Iowa
DNA Sequencing by: Dr. M. Bento Soares, University of Iowa
Clone Distribution: Researchers may obtain clones from Research
Genetics (www.resgen.com) or from Open Blosystems
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Bonaldo, M.P., Lennon, G. and Soares, M.B.
Normalization and subtraction: two approaches to facilitate gene
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bento-soares@miowa.edu
TAG INSURE-Lung Epithelial Cells Tissue nos 359-368
TAG LIBLUT-FF-ENU
TAG_EGGCTGTAGGC"
                                                                                                                                                                                                                                                                                                                                                                                                                                                                    University of Iowa
2024 University of Iowa Med Labs, Iowa City, IA 52242, USA
Tel: 319 356 4866
Fax: 319 356 7171
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Genome Res. 6 (9), 791-806 (1996)
97044477
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Seq primer: M13 FORWARD
POLYA=Yes.
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EST.
                                                                                                                                                                                                                        Homo sapiens (human)
Homo sapiens
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//db xref="texton:9606"
/cell line="HT1080"
/clone_lib="Athersys RAGE Library"
/clone_lib="Athersys RAGE Library"
/note="See" Creation of Gennee-wide Protein Expression
Libraries using Random Activation of Gene Expression
/ Nature Biotechnology, in press. Note that even though the
cell type indicated is HT1080, since a random activation
method was used, these sequence tags are not necessarily
expressed in HT1080 under normal circumstances."
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Pred. No. 7.1e-148;
0; Mismatches 15; Indels
                            /organism="Homo sapiens"
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